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=> d que 142
              1 SEA FILE=REGISTRY ABB=ON PLU=ON 7664-41-7/RN
L3
              1 SEA FILE=REGISTRY ABB=ON PLU=ON
L5
                                                  1314-62-1/RN
              1 SEA FILE=REGISTRY ABB=ON PLU=ON 7440-06-4/RN
1 SEA FILE=REGISTRY ABB=ON PLU=ON 11122-73-9/RN
L6
L7.
         227949 SEA FILE=HCAPLUS ABB=ON PLU=ON L3 OR AMMONIA
L8
          25987 SEA FILE=HCAPLUS ABB=ON PLU=ON L5 OR VANADIA
L9
L10
         347310 SEA FILE=HCAPLUS ABB=ON PLU=ON L6 OR PLATINUM OR PT
T-12
           2926 SEA FILE=HCAPLUS ABB=ON
                                         PLU=ON L7
L13
              1 SEA FILE=HCAPLUS ABB=ON
                                         PLU=ON
                                                 L8 AND L9 AND L10 AND L12
L14
            325 SEA FILE=HCAPLUS ABB=ON PLU=ON
                                                 "REFRACTORY METAL
                OXIDES"+PFT,NT/CT
             10 SEA FILE=HCAPLUS ABB=ON PLU=ON L14 AND L8
L15 .
             16 SEA FILE=HCAPLUS ABB=ON PLU=ON L8 AND REFRACTORY METAL
L16
                OXIDE?
             16 SEA FILE=HCAPLUS ABB=ON PLU=ON L15 OR L16
L17
            5 SEA FILE=HCAPLUS ABB=ON PLU=ON L17 AND L10
L18
             3 SEA FILE=HCAPLUS ABB=ON PLU=ON L18 AND L9
L19
             16 SEA FILE=HCAPLUS ABB=ON PLU=ON (L17 OR L18 OR L19)
L20
             16 SEA FILE=HCAPLUS ABB=ON PLU=ON L13 OR L20
L21
L22
             1 SEA FILE=HCAPLUS ABB=ON PLU=ON LAYERED AMMONIA OXIDAT?
                QUE ABB=ON PLU=ON FILM? OR THINFILM? OR LAYER? OR OVER
L25
              LAY? OR OVERLAID? OR LAMIN? OR LAMEL? OR MULTILAYER? OR S
                HEET? OR LEAF? OR FOIL? OR COAT? OR VENEER? OR SHEATH? OR
                 COVER?
L26
          20003 SEA FILE=HCAPLUS ABB=ON PLU=ON L8(L)L25
L28
             18 SEA FILE=HCAPLUS ABB=ON PLU=ON L26 AND L10 AND L9
L29
             15 SEA FILE=HCAPLUS ABB=ON PLU=ON L28 AND CAT/RL
L30
          91484 SEA FILE=HCAPLUS ABB=ON PLU=ON
                                                 "OXIDATION CATALYSTS"+PFT;
                NT/CT
L31
              5 SEA FILE=HCAPLUS ABB=ON PLU=ON L29 AND L30
L33
             29 SEA FILE=HCAPLUS ABB=ON PLU=ON L21 OR L22 OR L29 OR L31
L34
             17 SEA FILE=HCAPLUS ABB=ON PLU=ON L33 AND AIR POLLU?/SC,SX
             12 SEA FILE=HCAPLUS ABB=ON PLU=ON L33 NOT L34
L35
L36
              6 SEA FILE=HCAPLUS ABB=ON PLU=ON L35 AND CAT?
                QUE ABB=ON PLU=ON SUBSTRAT? OR SURFACE? OR BASE# OR SU
L37
                BSTRUCT? OR UNDERSTRUCT? OR UNDERLAY? OR FOUNDATION? OR P
                ANE? OR DISK? OR DISC# OR WAFER?
L38
            780 SEA FILE=HCAPLUS ABB=ON PLU=ON L37 AND L12
L39
              1 SEA FILE=HCAPLUS ABB=ON PLU=ON L38 AND L8 AND L9 AND L10
             3 SEA FILE=HCAPLUS ABB=ON PLU=ON L38 AND L8
L40
L41
             4 SEA FILE=HCAPLUS ABB=ON PLU=ON L36 AND L37
L42
             23 SEA FILE=HCAPLUS ABB=ON PLU=ON L34 OR (L39 OR L40 OR
              L41)
=> d que 156
                QUE ABB=ON PLU=ON SUBSTRAT? OR SURFACE? OR BASE# OR SU
L37
                BSTRUCT? OR UNDERSTRUCT? OR UNDERLAY? OR FOUNDATION? OR P
                ANE? OR DISK? OR DISC# OR WAFER?
            530 SEA FILE-WPIX ABB-ON PLU-ON REFRACTORY METAL OXIDE?
L44
             13 SEA FILE=WPIX ABB=ON PLU=ON L44 AND AMMONIA
L45
              4 SEA FILE=WPIX ABB=ON · PLU=ON L45 AND PLATINUM?
L46
              1 SEA FILE=WPIX ABB=ON PLU=ON L45 AND VANADIA?
L47
             4 SEA FILE-WPIX ABB-ON PLU-ON L46 OR L47
L48
            3 SEA FILE-WPIX ABB-ON PLU-ON L45 AND B01D0053?/IPC 5 SEA FILE-WPIX ABB-ON PLU-ON L48 OR L49
L49
L50
L51
            67 SEA FILE=WPIX ABB=ON PLU=ON
                                             L44 AND PLATINUM
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33 SEA FILE=WPIX ABB=ON PLU=ON L51 AND B01D0053?/IPC
17 SEA FILE=WPIX ABB=ON PLU=ON L52 AND L37
17 SEA FILE=WPIX ABB=ON PLU=ON L53 AND CATALYST?
2 SEA FILE=WPIX ABB=ON PLU=ON/ L54 AND (AMMONIA OR NH3)
L52
L53
L54
L55
              5 SEA FILE=WPIX ABB=ON PLU=ON L50 OR L55
L56
=> d que 157
      530 SEA FILE=WPIX ABB=ON PLU=ON REFRACTORY METAL OXIDE?
L44
             O SEA FILE=COMPENDEX ABB=ON PLU=ON L44 AND AMMONIA
L57
=> d que 161
             530 SEA FILE=WPIX ABB=ON PLU=ON REFRACTORY METAL OXIDE?
              O SEA FILE=PASCAL ABB=ON PLU=ON L44 AND AMMONIA
             14 SEA FILE=PASCAL ABB=ON PLU=ON REFRACTORY METAL OXIDE?
             O SEA FILE=PASCAL ABB=ON PLU=ON L59 AND (AMMONIA OR NH3)
L60
             O SEA FILE=PASCAL ABB=ON PLU=ON L58 OR L60
L61
=> d que 167
             530 SEA FILE-WPIX ABB-ON PLU-ON REFRACTORY METAL OXIDE?
L62
              O SEA FILE=JAPIO ABB=ON PLU=ON L44 AND AMMONIA
              27 SEA FILE=JAPIO ABB=ON PLU=ON REFRACTORY METAL OXIDE?
L63
              O SEA FILE=JAPIO ABB=ON PLU=ON L63 AND NH3
L64
              4 SEA FILE=JAPIO ABB=ON PLU=ON L63 AND PLATINUM
L65
             O SEA FILE=JAPIO ABB=ON PLU=ON L63 AND VANADIA
L66
             4 SEA FILE=JAPIO ABB=ON PLU=ON L62 OR (L64 OR L65 OR L66)
L67
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=> dup rem 142 156 157 161 167
L57 HAS NO ANSWERS
L61 HAS NO ANSWERS
FILE 'HCAPLUS' ENTERED AT 11:45:50 ON 28 AUG 2007
USE IS SUBJECT TO THE TERMS OF YOUR STN CUSTOMER AGREEMENT.
PLEASE SEE "HELP USAGETERMS" FOR DETAILS.
COPYRIGHT (C) 2007 AMERICAN CHEMICAL SOCIETY (ACS)

FILE 'WPIX' ENTERED AT 11:45:50 ON 28 AUG 2007 COPYRIGHT (C) 2007 THE THOMSON CORPORATION

FILE 'JAPIO' ENTERED AT 11:45:50 ON 28 AUG 2007

COPYRIGHT (C) 2007 Japanese Patent Office (JPO) - JAPIO

PROCESSING COMPLETED FOR L42

PROCESSING COMPLETED FOR L56

PROCESSING COMPLETED FOR L61

PROCESSING COMPLETED FOR L61

PROCESSING COMPLETED FOR L67

L68 29 DUP REM L42 L56 L57 L61 L67 (3 DUPLICATES REMOVED)

ANSWERS '1-23' FROM FILE HCAPLUS

ANSWERS '24-25' FROM FILE WPIX

ANSWERS '26-29' FROM FILE JAPIO

=> d 1-23 ibib ed abs hitstr hitind

L68 ANSWER 1 OF 29 HCAPLUS COPYRIGHT 2007 ACS on STN DUPLICATE 1
ACCESSION NUMBER: 2006:168226 HCAPLUS
DOCUMENT NUMBER: 144:217801
TITLE: Zone coated catalyst to simultaneously reduce NOx and unreacted ammonia

```
INVENTOR(S):
```

PATENT ASSIGNEE(S):

SOURCE:

Patchett, Joseph Allan; Dettling, Joseph Charles Engelhard Corporation, USA U.S. Pat. Appl. Publ., 22 pp.

CODEN: USXXCO

DOCUMENT TYPE: LANGUAGE:

Patent English

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

PA'	PATENT NO.			KIN	D	DATE					ION 1			D	ATE	_	
ับร	2006	0398	43		A1	-	2006	0223							2	004082	3
WO	2006	0239:	32		A1		2006	0302	1	WO 2	005-1	US29:	992		2	005082	2
	W:	ΑE,	AG,	AL,	AM,	ΑT,	AU,	ΑZ,	BA,	BB,	BG,	BR,	BW,	BY,	ΒZ,	CA,	
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		GB,	GD,	GE,	GH,	GM,	HR,	HU,	ID,	IL,	IN,	IS,	JP,	KE,	KG,	KM,	
		ΚP,	KR,	KZ,	LC,	LK,	LR,	LS,	LT,	LU,	LV,	MA,	MD,	MG,	MK,	MN,	
		MW,	MX,	MZ,	NA,	NG,	NI,	NO,	NZ,	OM,	PG,	PH,	PL,	PT,	RO,	RU,	
		SC,	SD,	SE,	SG,	SK,	SL,	SM,	SY,	ТJ,	TM,	TN,	TR,	TT,	TZ,	UA,	
		UG,	US,	UΖ,	VC,	VN,	ΥU,	ZA,	ZM,	ZW							
	RW:	ΑT,	BE,	BG,	CH,	CY,	CZ,	DE,	DK,	EE,	ES,	FI,	FR,	GB,	GR,	HU,	
		ΙE,	IS,	IT,	LT,	LU,	ĽV,	MC,	NL,	PL,	PT,	RO,	SE,	SI,	SK,	TR,	
		BF,	ВJ,	CF,	CG,	CI,	CM,	GΑ,	GN,	GQ,	GW,	ML,	MR,	ΝE,	SN,	TD,	
							LS,										
		ZW,	AM,	ΑZ,	BY,	KG,	KZ,	MD,	RU,	TJ,	TM						
EP	1784	258			A1		2007	0516		EP 20	005-	79394	42		2	005082	2
	R:	AT,	BE,	BG,	CH,	CY,	CZ,	DE,	DK,	EE,	ES,	FI,	FR,	GB,	GR,	HU,	
		IE,	IS,	IT,	LI,	LT,	LU,	LV,	MC,	NL,	PL,	PT,	RO,	SE,	SI,	SK, TI	R
PRIORIT	Y APP															004082	

Entered STN: 23 Feb 2006 ED

AB Provided is an emissions treatment system and method for reducing NOx emissions in the exhaust stream produced from an internal combustion engine. The system has an injector for periodically metering ammonia or an ammonia precursor into an exhaust stream; and a first substrate with a first selective catalytic reduction (SCR) catalyst composition, downstream of the injector. The first substrate has an inlet end, an outlet end, a length extending between the inlet end to the outlet end, wall elements and a plurality of passages defined by the wall elements. The first SCR catalyst composition is disposed on the wall elements from the inlet end toward the outlet end to a length that is less than the substrate's axial length to form an inlet zone. The first substrate also has an NH3 destruction catalyst composition with a platinum group metal component dispersed on a refractory metal oxide. The NH3 destruction catalyst is disposed on the wall elements from the outlet end toward the inlet end to a length that is less than the substrate's axial length to form an outlet zone. Generally, there is from 0.1 to 10 g/ft3 of platinum group metal component in the outlet zone.

WO 2005-US29992

20050822

1314-62-1, Vanadia, uses

(as component of first SCR catalyst composition; zone coated catalyst to simultaneously reduce NOx and unreacted ammonia

RN 1314-62-1 HCAPLUS

CN Vanadium oxide (V2O5) (CA INDEX NAME)

*** STRUCTURE DIAGRAM IS NOT AVAILABLE ***

```
IT
     7664-41-7, Ammonia, processes
        (slip, reduction of; zone coated catalyst to simultaneously
        reduce NOx and unreacted ammonia)
RN
     7664-41-7 HCAPLUS
     Ammonia (CA INDEX NAME)
CN
NH3
IT
     7440-06-4, Platinum, uses
        (zone coated catalyst to simultaneously reduce NOx and
        unreacted ammonia)
RN
     7440-06-4 HCAPLUS
     Platinum (CA INDEX NAME)
CN
Pt
INCL 423239100; 422177000; 422180000; 422172000
     59-3 (Air Pollution and Industrial Hygiene)
     Section cross-reference(s): 67
ST
     zone coated catalyst SCR nitrogen oxide removal
     ammonia slip
IT
     Zeolites (synthetic), uses
        (Cu- or Fe-exchanged, as component of first SCR catalyst composition;
        zone coated catalyst to simultaneously reduce NOx and
        unreacted ammonia)
IT
     Reduction
        (selective catalytic; zone coated catalyst to
        simultaneously reduce NOx and unreacted ammonia)
IT
     Exhaust gases (engine)
        (zone coated catalyst to simultaneously reduce NOx and
        unreacted ammonia)
IT
     Platinum-group metals
        (zone coated catalyst to simultaneously reduce NOx and
        unreacted ammonia)
IT
     1306-38-3, Ceria, uses
        (as catalyst for NH3 destruction; zone coated catalyst to
        simultaneously reduce NOx and unreacted ammonia)
IT
     1314-35-8, Tungsten oxide (WO3), uses 1314-62-1,
     Vanadia, uses
                   13463-67-7, Titania, uses
        (as component of first SCR catalyst composition; zone coated
        catalyst to simultaneously reduce NOx and unreacted ammonia
IT
     7664-41-7, Ammonia, processes
        (slip, reduction of; zone coated catalyst to simultaneously
        reduce NOx and unreacted ammonia)
IT
     7439-88-5, Iridium, uses 7440-05-3, Palladium, uses
     7440-06-4, Platinum, uses 7440-16-6, Rhodium, uses
        (zone coated catalyst to simultaneously reduce NOx and
        unreacted ammonia)
IT
     11104-93-1, Nitrogen oxide, processes
        (zone coated catalyst to simultaneously reduce NOx and
        unreacted ammonia)
L68 ANSWER 2 OF 29 HCAPLUS COPYRIGHT 2007 ACS on STN DUPLICATE 2
ACCESSION NUMBER:
                         2005:451605 HCAPLUS
```

142:468240

DOCUMENT NUMBER:

```
Emissions treatment system with NSR and SCR
TITLE:
                         catalysts
INVENTOR (S):
                         Li, Yuejin; Deeba, Michel; Dettling, Joseph
                         Charles; Patchett, Joseph Allan; Roth, Stanley
                         Allan
PATENT ASSIGNEE(S):
                         Engelhard Corporation, USA
SOURCE:
                         PCT Int. Appl., 44 pp.
                         CODEN: PIXXD2
DOCUMENT TYPE:
                         Patent
LANGUAGE:
                         English
FAMILY ACC. NUM. COUNT:
PATENT INFORMATION:
     PATENT NO.
                         KIND
                                DATE
                                            APPLICATION NO.
                                                                   DATE
     -----
                         ----
                                -----
                                            -----
                                                                   -----
     WO 2005047663
                          A2
                                20050526
                                            WO 2004-US36723
                                                                   20041104
     WO 2005047663
                         Α3
                                20050623
         W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA,
             CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI,
             GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP,
             KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW,
             MX, MZ, NA, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD,
             SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ,
             VC, VN, YU, ZA, ZM, ZW
         RW: BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW,
            AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ,
            DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LU, MC, NL,
             PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN,
            GQ, GW, ML, MR, NE, SN, TD, TG
     US 2005129601
                                20050616
                                          US 2004-975428
                         A1
                                                                   20041029
                                20060809
                                            EP 2004-800722
     EP 1687514
                          A2
                                                                   20041104
            AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC,
             PT, IE, SI, FI, RO, CY, TR, BG, CZ, EE, HU, PL, SK, IS
     IN 2006KN01164
                         .A
                                20070427
                                            IN 2006-KN1164
                                                                   20060504
PRIORITY APPLN. INFO.:
                                            US 2003-517137P
                                                                   20031104
                                            US 2004-975428
                                                                   20041029
                                            WO 2004-US36723
                                                                   20041104
ED
    Entered STN:
                  27 May 2005
     Provided is an emissions treatment system for an exhaust stream,
AB
     having a NOx storage reduction (NSR) catalyst with a NOx sorbent at a
     concentration of at least 0.1g/in3 and a platinum group metal
     component dispersed on a refractory metal
     oxide support; and a selective catalytic reduction (SCR) catalyst
     disposed downstream of the NSR catalyst. The emissions treatment.
     system is advantageously used for the treatment of exhaust streams
     from diesel engines and lean burn gasoline engines.
IT
     7440-06-4, Platinum, uses
        (NSR catalyst component; engine exhaust treatment system with NOx
        storage reduction and selective catalytic reduction catalysts)
     7440-06-4 HCAPLUS
RN
     Platinum (CA INDEX NAME)
CN
```

Pt

IT 1314-62-1, Vanadium pentoxide, uses

(SCR catalyst component; engine exhaust treatment system with NOx storage reduction and selective catalytic reduction catalysts)

1314-62-1 HCAPLUS RN

CN Vanadium oxide (V2O5) (CA INDEX NAME)

*** STRUCTURE DIAGRAM IS NOT AVAILABLE ***

7664-41-7, Ammonia, reactions IT

(SCR reductant; engine exhaust treatment system with NOx storage reduction and selective catalytic reduction catalysts)

RN 7664-41-7 HCAPLUS

Ammonia (CA INDEX NAME) CN

NH3

ICM F01N003-08 IC

ICS F01N003-20

CC 59-3 (Air Pollution and Industrial Hygiene) Section cross-reference(s): 67

IT Refractory metal oxides

> (substrate for NOx sorbent component of NSR catalyst; engine exhaust treatment system with NOx storage reduction and selective catalytic reduction catalysts)

IT 7440-05-3, Palladium, uses 7440-06-4, Platinum,

7440-16-6, Rhodium, uses

(NSR catalyst component; engine exhaust treatment system with NOx storage reduction and selective catalytic reduction catalysts)

IT 1314-35-8, Tungsten oxide (WO3), uses 1314-62-1, Vanadium pentoxide, uses 13463-67-7, Titania, uses

> (SCR catalyst component; engine exhaust treatment system with NOx storage reduction and selective catalytic reduction catalysts)

IT 7664-41-7, Ammonia, reactions

(SCR reductant; engine exhaust treatment system with NOx storage reduction and selective catalytic reduction catalysts)

ANSWER 3 OF 29 HCAPLUS COPYRIGHT 2007 ACS on STN DUPLICATE 3

ACCESSION NUMBER:

2005:220099 HCAPLUS

DOCUMENT NUMBER:

142:265825

TITLE:

Layered ammonia oxidation catalyst

INVENTOR(S):

Tran, Pascaline Harrison; Chen, James Mon-Her; Lapadula, Gerard Diomede; Blute, Marc Thomas

PATENT ASSIGNEE(S):

SOURCE:

Engelhard Corporation, USA

U.S. Pat. Appl. Publ., 5 pp. CODEN: USXXCO

DOCUMENT TYPE:

Patent

LANGUAGE:

English 1

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

PATENT NO.	KIND DATE	APPLICATION NO.	DATE
US 2005054524	A1 20050310	US 2003-659159	20030910
WO 2005025724	A1 20050324	WO 2004-US27717	20040826
W: AE, AG, AL,	AM, AT, AU, AZ,	BA, BB, BG, BR, BW, BY	, BZ, CA,
CH, CN, CO,	CR, CU, CZ, DE,	DK, DM, DZ, EC, EE, EG	, ES, FI,
GB, GD, GE,	GH, GM, HR, HU,	ID, IL, IN, IS, JP, KE	, KG, KP,
KR, KZ, LC,	LK, LR, LS, LT,	LU, LV, MA, MD, MG, MK	, MN, MW,
MX, MZ, NA,	NI, NO, NZ, OM,	PG, PH, PL, PT, RO, RU	, SC, SD,

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SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ,
         VC, VN, YU, ZA, ZM, ZW
RW: BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW,
             AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ,
             DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PL,
             PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ,
             GW, ML, MR, NE, SN, TD, TG
     EP 1660216
                          A1
                                 20060531
                                             EP 2004-782238
                                                                     20040826
         R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC,
             PT, IE, SI, FI, RO, CY, TR, BG, CZ, EE, HU, PL, SK
                                 20061018
                                             CN 2004-80026110
                                                                     20040826
     CN 1849163
                          Α
     JP 2007504945
                          Т
                                 20070308
                                             JP 2006-526126
                                                                     20040826
                                             US 2003-659159
PRIORITY APPLN. INFO.:
                                                                     20030910
                                             WO 2004-US27717
                                                                     20040826
ED
     Entered STN: 13 Mar 2005
AB
     The invention pertains to a layered ammonia
     oxidation catalyst. The layered catalyst causes
     ammonia to be selectively oxidized in the presence of an
     oxidant such as air, while minimizing the formation of nitrogen oxides
     (NOx). The layered catalyst comprises a refractory oxide
     support such as gamma alumina upon which a platinum
     component is deposited and a vanadia component is deposited
     on the platinum. The catalyst is preferably disposed on a
     substrate such as a metal foil whose surface
     contains a "herringbone" pattern.
TT
     1314-62-1, Vanadia, processes 7440-06-4,
     Platinum, processes 11122-73-9
        (layered ammonia oxidation catalyst)
RN
     1314-62-1 HCAPLUS
CN
     Vanadium oxide (V2O5)
                            (CA INDEX NAME)
    STRUCTURE DIAGRAM IS NOT AVAILABLE ***
RN
     7440-06-4 HCAPLUS
CN
     Platinum
              (CA INDEX NAME)
Pt
RN
     11122-73-9 HCAPLUS
CN
     Chromium alloy, nonbase, Cr, Fe (CA INDEX NAME)
             Component
Component
          Registry Number
=======+===+=========
               7440-47-3
    Cr .
               7439-89-6
    Fe
IT
     7664-41-7, Ammonia, reactions
        (layered ammonia oxidation catalyst)
     7664-41-7 HCAPLUS
RN.
     Ammonia (CA INDEX NAME)
CN
```

NH3

IC ICM B01J023-648

```
INCL 502312000; 423237000
     59-4 (Air Pollution and Industrial Hygiene)
     Section cross-reference(s): 67
ST
     layered ammonia oxidn catalyst
     platinum vanadia alumina
     Air pollution
IT
        (control; layered ammonia oxidation
        catalyst)
ΙT
     Combustion gases
     Flue gases
     Honeycomb structures
       Oxidation catalysts
       Surface area
     Waste gases
        (layered ammonia oxidation catalyst)
IT
     Refractory metal oxides
        (layered ammonia oxidation catalyst)
IT
     1344-28-1, Alumina, processes
        (gamma; layered ammonia oxidation
        catalyst)
IT
     1314-62-1, Vanadia, processes 7440-06-4,
     Platinum, processes 11122-73-9
        (layered ammonia oxidation catalyst)
IT
     7664-41-7, Ammonia, reactions
        (layered ammonia oxidation catalyst)
L68 ANSWER 4 OF 29 HCAPLUS COPYRIGHT 2007 ACS on STN
ACCESSION NUMBER:
                         2005:138868 HCAPLUS
DOCUMENT NUMBER:
                         142:224506
                         Catalyst arrangement and method of purifying the
TITLE:
                         exhaust gas of internal combustion engines
                         operated under lean conditions
INVENTOR(S):
                         Pfeifer, Marcus; Soeger, Nicola; Demel, Yvonne;
                         Kuhl, Tobias; Spurk, Paul Christian; Gieshoff,
                         Juergen; Lox, Egbert; Kreuzer, Thomas
PATENT ASSIGNEE(S):
                         Umicore A.-G. & Co. K.-G., Germany
                         PCT Int. Appl., 19 pp.
SOURCE:
                         CODEN: PIXXD2
DOCUMENT TYPE:
                         Patent
LANGUAGE:
                         English
FAMILY ACC. NUM. COUNT:
                         1
PATENT INFORMATION:
```

PAT	CENT :	NO.			KIN	נ כ	DATE		1	APPL	ICAT:	ION	NO.		D	ATE
WO	2005	0141	 46		A1	- · :	2005	0217	1	WO 2	 004-1	 EP85:	39		2	0040729
	W:	AE,	AG,	AL,	AM,	AT,	AU,	ΑZ,	BA,	BB,	BG,	BR,	BW,	BY,	BZ,	CA,
		CH,	CN,	CO,	CR,	CU,	CZ,	DE,	DK,	DM,	DZ,	EC,	EE,	EG,	ES,	FI,
		GB,	GD,	GE,	GH,	GM,	HR,	HU,	ID,	IL,	IN,	IS,	JP,	ΚE,	KG,	KP,
•		KR,	ΚZ,	LC,	LK,	LR,	LS,	LT,	LU,	LV,	MA,	MD,	.MG,	MK,	MN,	MW,
	•	MX,	ΜZ,	NA,	NI,	NO,	NZ,	OM,	PG,	PH,	PL,	PT,	RO,	RU,	SC,	SD,
		SE,	SG,	SK,	SL,	SY,	ТJ,	TM,	TN,	TR,	TT,	TZ,	UA,	ŪĠ,	US,	UZ,
		VC,	VN,	ΥU,	ZA,	ZM,	ZW									
	RW:	BW,	GH,	GM,	KE,	LS,	MW,	MZ,	NA,	SD,	SL,	SZ,	TZ,	UG,	ZM,	ZW,
		AM,	ΑZ,	BY,	KG,	ΚZ,	MD,	RU,	TJ,	TM,	AT,	BE,	BG,	CH,	CY,	CZ,
		DE,	DK,	EE,	ES,	FI,	FR,	GB,	GR,	HU,	IE,	IT,	LU,	MC,	NL,	PL,
		PT,	RO,	SE,	SI,	SK,	TR,	BF,	ВJ,	CF,	CG,	CI,	CM,	GΑ,	GN,	GQ,
		GW,	ML.,	MR,	ΝE,	SN,	TD,	TG								
DE	1033	5785			A1	:	2005	0310]	DE 20	003-	1033	5785		2	0030805
CA	2534	806			A1	:	20.05	0217	(CA 2	004-	2534	806		20	0040729

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EP 1660217
                              A1
                                     20060531
                                                   EP 2004-763630
                                                                             20040729
              AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, FI, RO, CY, TR, BG, CZ, EE, HU, PL, SK
     BR 2004013367
                             Α
                                     20061017
                                                  BR 2004-13367
                                                                             20040729
                                                                             20040729
                             Α
                                     20061115
                                                   CN 2004-80029120
     CN 1863586
                             Т
                                     20070125
                                                  JP 2006-522295
     JP 2007501107
                                                                             20040729
                             A1
                                     20070517
                                                  US 2006-567204
     US 2007110650
                                                                             20061211
PRIORITY APPLN. INFO.:
                                                  DE 2003-10335785
                                                                          A 20030805
                                                  WO 2004-EP8539
                                                                          W 20040729
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ED Entered STN: 17 Feb 2005

AB The invention relates to a catalyst arrangement for purifying the exhaust gases of internal combustion engines operated under lean conditions. It is proposed that a thin-walled, porous carrier be coated on the exit surface by a catalyst for selective catalytic reduction and on the entry surface by a catalyst layer able to store nitrogen oxides under lean exhaust gas conditions and to convert nitrogen oxides into ammonia under rich exhaust gas conditions. When the exhaust gas is passed through the catalytic coatings and the support material, a significant improvement in the nitrogen oxide conversion is achieved compared to a series arrangement of the catalysts on sep. carriers. Wall flow filters have been found to be useful as thin-walled carriers.

IT 7440-06-4, Platinum, uses

(NOx storage catalyst; lean burn engine exhaust treatment using thin-walled, porous carrier coated on entry surface with NOx storage catalyst and on exit surface with selective reduction catalyst)

RN 7440-06-4 HCAPLUS

CN Platinum (CA INDEX NAME)

Pt

IT 1314-62-1, Vanadium oxide (V2O5), uses

(SCR catalyst component; lean burn engine exhaust treatment using thin-walled, porous carrier coated on entry surface with NOx storage catalyst and on exit surface with selective reduction catalyst)

RN 1314-62-1 HCAPLUS

CN Vanadium oxide (V2O5) (CA INDEX NAME)

*** STRUCTURE DIAGRAM IS NOT AVAILABLE ***

IC ICM B01D053-94

ICS F01N003-28

CC 59-3 (Air Pollution and Industrial Hygiene)
Section cross-reference(s): 67

IT Alkaline earth oxides

Platinum-group metals

(NOx storage catalyst; lean burn engine exhaust treatment using thin-walled, porous carrier coated on entry surface with NOx storage catalyst and on exit surface with selective reduction catalyst)

IT 513-77-9, Barium carbonate 7440-06-4, Platinum,

uses 7440-16-6, Rhodium, uses 65453-23-8, Cerium zirconium oxide (NOx storage catalyst; lean burn engine exhaust treatment using thin-walled, porous carrier coated on entry surface with NOx storage catalyst and on exit surface with selective reduction catalyst)

IT 1313-27-5, Molybdenum oxide, uses 1314-35-8, Tungsten oxide (WO3),
 uses 1314-62-1, Vanadium oxide (V2O5), uses 7631-86-9,
 Silica, uses 13463-67-7, Titania, uses

(SCR catalyst component; lean burn engine exhaust treatment using thin-walled, porous carrier coated on entry surface with NOx

storage catalyst and on exit surface with selective reduction catalyst) REFERENCE COUNT: THERE ARE 7 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT L68 ANSWER 5 OF 29 HCAPLUS COPYRIGHT 2007 ACS on STN ACCESSION NUMBER: 2006:151542 HCAPLUS DOCUMENT NUMBER: 144:455364 TITLE: Pt-V2O5-WO3/TiO2 catalysts supported on SiC filter for NO reduction at low temperature AUTHOR (S): Choi; Joo-Hong; Kim, Jin-Hyun; Bak, Young-Cheoul; Amal, Rose; Scott, Jason CORPORATE SOURCE: Department of Chemical and Biological Engineering/ERI, Gyeongsang National University, Jinju, 660-701, S. Korea SOURCE: Korean Journal of Chemical Engineering (2005), 22(6), 844-851 CODEN: KJCHE6; ISSN: 0256-1115 PUBLISHER: Korean Institute of Chemical Engineers DOCUMENT TYPE: Journal LANGUAGE: English ED Entered STN: 17 Feb 2006 AB A catalytic filter, V2O5-WO3-TiO2 supported on a ceramic filter, is a promising material to simultaneously treat particulates and NOx at optimum temps. of .apprx.320°. To improve its catalytic activity at low temps., the effect of Pt addition on the catalytic filter was studied. Pt-V2O5-WO3-TiO2/SiC filters were prepared by co-impregnation of Pt, V, and W precursors on a TiO2 coated-SiC filter by vacuum aided-dip coating. The Pt -added catalytic filter shifted optimum working temps. from 280-330° (for non-Pt-impregnated filter) to 180-230°, providing a Nx slip concentration <20 ppm for treating 700 ppm NO at a face velocity of 2 cm/s with the same value over the non-Pt-added catalytic filter. The promotional effect following Pt addition is believed to result from elec. modification of the catalyst maintaining a high electron transfer state. NH3 oxidation was also observed to be dominant above the optimal selective catalytic reduction temperature IT 7664-41-7, Ammonia, reactions (reductant; low temperature selective catalytic reduction of waste gas nitric oxide by ammonia over platinum-promoted vanadia/tungsten oxide titania-coated silicon carbide catalyst) RN7664-41-7 HCAPLUS CN Ammonia (CA INDEX NAME) NH3 IT 1314-62-1, Vanadia, uses

-promoted tungsten oxide and; low temperature selective catalytic reduction

(titania-coated silicon carbide, platinum

of waste gas nitric oxide by ammonia over

coated silicon carbide catalyst)

1314-62-1 HCAPLUS

RN

platinum-promoted vanadia/tungsten oxide titania-

```
Vanadium oxide (V2O5)
                            (CA INDEX NAME)
CN
*** STRUCTURE DIAGRAM IS NOT AVAILABLE ***
IT
     7440-06-4, Platinum, uses
        (titania-coated silicon carbide, vanadia
        /tungsten oxide doped with; low temperature selective catalytic reduction of
        waste gas nitric oxide by ammonia over platinum
        -promoted vanadia/tungsten oxide titania-coated
        silicon carbide catalyst)
     7440-06-4 HCAPLUS
RN
CN
     Platinum (CA INDEX NAME)
Pt
CC
     59-4 (Air Pollution and Industrial Hygiene)
     Section cross-reference(s): 57, 67
ST
     silicon carbide supported platinum vanadia
     tungsten titania redn catalyst; ammonia selective catalytic redn waste
     gas nitric oxide
IT
     Reduction catalysts
        (Pt-V2O5-WO3/TiO2-SiC; low temperature selective catalytic
        reduction of waste gas nitric oxide by ammonia over
        platinum-promoted vanadia/tungsten oxide titania-
        coated silicon carbide catalyst)
IT
     Flue gases
     Waste gases
        (low temperature selective catalytic reduction of waste gas nitric oxide by
        ammonia over platinum-promoted vanadia
        /tungsten oxide titania-coated silicon carbide catalyst)
IT
     7727-37-9, Nitrogen, processes
                                      10024-97-2, Nitrous oxide, processes
     10102-44-0, Nitrogen dioxide, processes
        (low temperature selective catalytic reduction of waste gas nitric oxide by
        ammonia over platinum-promoted vanadia
        /tungsten oxide titania-coated silicon carbide catalyst)
IT
     7664-41-7, Ammonia, reactions
        (reductant; low temperature selective catalytic reduction of waste gas
nitric
        oxide by ammonia over platinum-promoted
       vanadia/tungsten oxide titania-coated silicon
        carbide catalyst)
     13463-67-7, Titania, uses
IT
        (silicon carbide coated with; low temperature selective
        catalytic reduction of waste gas nitric oxide by ammonia over
       platinum-promoted vanadia/tungsten oxide titania-
        coated silicon carbide catalyst)
    1314-62-1, Vanadia, uses
        (titania-coated silicon carbide, platinum
        -promoted tungsten oxide and; low temperature selective catalytic reduction
       of waste gas nitric oxide by ammonia over
       platinum-promoted vanadia/tungsten oxide titania-
        coated silicon carbide catalyst)
IT
     1314-35-8, Tungsten oxide, uses
        (titania-coated silicon carbide, platinum
        -promoted vanadia and; low temperature selective catalytic
        reduction of waste gas nitric oxide by ammonia over
       platinum-promoted vanadia/tungsten oxide titania-
        coated silicon carbide catalyst)
IT
    7440-06-4, Platinum, uses
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(titania-coated silicon carbide, vanadia
/tungsten oxide doped with; low temperature selective catalytic reduction of
waste gas nitric oxide by ammonia over platinum
-promoted vanadia/tungsten oxide titania-coated
silicon carbide catalyst)

IT 409-21-2, Silicon carbide, uses

(titania-coated, platinum-promoted

vanadia/tungsten oxide; low temperature selective catalytic reduction
of waste gas nitric oxide by ammonia over

platinum-promoted vanadia/tungsten oxide titania-

coated silicon carbide catalyst)

REFERENCE COUNT:

THERE ARE 38 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE

RE FORMAT

L68 ANSWER 6 OF 29 HCAPLUS COPYRIGHT 2007 ACS on STN

38 .

ACCESSION NUMBER:

2003:950220 HCAPLUS

DOCUMENT NUMBER:

139:397779

TITLE:

Pollutant reductions in engine exhaust gases by combustion of fuel emulsions and oxidation of exhaust gas components in flow-through oxidation

catalysts

INVENTOR(S):

Brown, Kevin F.; Langer, Deborah A.; Duncan, David

Α.

PATENT ASSIGNEE(S):

The Lubrizol Corporation, Can.

SOURCE:

U.S. Pat. Appl. Publ., 17 pp., Cont.-in-part of

U.S. Ser. No. 557,953.

CODEN: USXXCO

DOCUMENT TYPE:

Patent

LANGUAGE:

English

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
			·	
US 2003221360	A1	20031204	US 2003-457510	20030609
US 6949235	B2	20050927		
US 6606856	B1	20030819	US 2000-557953	20000424
PRIORITY APPLN. INFO.:			US 2000-519056	32 20000303
			IIS 2000-557953	2 20000424

ED Entered STN: 07 Dec 2003

AB Reduction of exhaust pollutants from engines, especially NOx, N2O, and particulates, is carried out by: (1) combusting an aqueous fuel emulsion that contains suitable emulsifying additives and combustion improvers, and (2) passing the exhaust gas from the engine into a flow-through cellular monolith containing an oxidation catalyst. The fuel emulsions, especially

diesel fuels, contain: (1) 1 or 2 fuel-soluble additives prepared by reacting C50-500-hydrocarbyl-substituted carboxylic acids (with different mol. wts.) with ammonia or an amine, (2) an ionic or nonionic surfactant with a hydrophilic-lipophilic balance of 1-40, (3) emulsion-stabilizing and combustion-improving water-soluble compds., such as amine or ammonium salts, azides, nitro compds., and alkali metal and alkaline earth metal salts, and (4) cetane improvers, antifreeze agents, and organic solvents. A metal or ceramic monolith coated with a washcoat material selected from zeolites, Al2O3, SiO2, TiO2, CeO2, ZrO2, V2O5, La2O3, and a catalyst selected from Pt, Pd, Rh, Ir, Ru, V, Cr, Mn, Fe, Co, Ni, Cu, Zn, Ag, Ce, and Ga.

1314-62-1, Vanadium oxide (V2O5), uses IT (washcoat catalyst support; pollutant redns. in engine exhaust gases by combustion of fuel emulsions and oxidation of exhaust gas components in flow-through oxidation catalysts)

1314-62-1 HCAPLUS RN

Vanadium oxide (V2O5) (CA INDEX NAME) CN

*** STRUCTURE DIAGRAM IS NOT AVAILABLE ***

7440-06-4, **Platinum**, uses ΙT

(washcoat oxidation catalyst; pollutant redns. in engine exhaust gases by combustion of fuel emulsions and oxidation of exhaust gas components in flow-through oxidation catalysts)

RN 7440-06-4 HCAPLUS

Platinum (CA INDEX NAME) CN

Pt

ICM C10L001-32

INCL 044301000

51-9 (Fossil Fuels, Derivatives, and Related Products) Section cross-reference(s): 59

IT Oxidation catalysts

(as washcoat on ceramic or metal monoliths; pollutant redns. in engine exhaust gases by combustion of fuel emulsions and oxidation of exhaust gas components in flow-through oxidation catalysts)

1306-38-3, Cerium oxide (CeO2), uses 1312-81-8, Lanthanum oxide TΤ 1314-23-4, Zirconium oxide (ZrO2), uses 1314-62-1, Vanadium oxide (V2O5), uses 1344-28-1, Alumina, uses 7631-86-9, 13463-67-7, Titania, uses Silica, uses

(washcoat catalyst support; pollutant redns. in engine exhaust gases by combustion of fuel emulsions and oxidation of exhaust gas components in flow-through oxidation catalysts)

7440-05-3, Palladium, uses IT 7439-88-5, Iridium, uses 7440-06-4, Platinum, uses 7440-16-6, Rhodium, uses

(washcoat oxidation catalyst; pollutant redns. in engine exhaust gases by combustion of fuel emulsions and oxidation of exhaust gas components in flow-through oxidation catalysts)

REFERENCE COUNT:

THERE ARE 64 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L68 ANSWER 7 OF 29 HCAPLUS COPYRIGHT 2007 ACS on STN

64

ACCESSION NUMBER:

2002:888632 HCAPLUS

DOCUMENT NUMBER:

137:374280

TITLE:

Catalyst for purification of diesel engine exhaust

gas

INVENTOR (S):

Kim, Young-Nam

PATENT ASSIGNEE(S):

KH Chemicals Co., Ltd., S. Korea

PCT Int. Appl., 58 pp. SOURCE:

CODEN: PIXXD2

DOCUMENT TYPE:

Patent

LANGUAGE:

English

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2002092224	. A1	20021121	WO 2001-KR845	20010522

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W:
              AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH,
              CN, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL,
              PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA,
              UG, US, UZ, VN, YU, ZA, ZW
          RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH,
              CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE,
              TR, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG
     KR 2002088013
                             Α
                                    20021125
                                                 KR 2001-26597
                                                                            20010516
     CA 2447665
                             A1
                                                 CA 2001-2447665
                                    20021121
                                                                            20010522
     AU 2001260735
                                                 AU 2001-260735
                             A1
                                    20021125
                                                                            20010522
     AU 2001260735
                             B2
                                    20070215
     BR 2001017021
                             Α
                                    20040420
                                                 BR 2001-17021
                                                                            20010522
     JP 2004513771
                             Т
                                    20040513
                                                 JP 2002-542591
                                                                            20010522
                             B2
     JP 3569703
                                    20040929
     CN 1524014
                             Α
                                    20040825
                                                 CN 2001-823429
                                                                            20010522
     US 2003104932
                             A1
                                    20030605
                                                 US 2002-958069
                                                                            20020716
     US 6855661
                            B2
                                    20050215
     TW 260241
                            В
                                    20060821
                                                 TW 2002-91133554
                                                                            20021115
     JP 2004105964
                            Α
                                    20040408
                                                 JP 2003-353417
                                                                            20031014
     MX 2003PA10426
                            Α
                                    20050921
                                                 MX 2003-PA10426
                                                                            20031114
     IN 2003MN01139
                            Α
                                    20050429
                                                 IN 2003-MN1139
                                                                            20031215
     US 2005032637
                             A1
                                    20050210
                                                 US 2004-936091
                                                                            20040907
PRIORITY APPLN. INFO.:
                                                 KR 2001-26597
                                                                        A 20010516
                                                 JP 2002-542591
                                                                        A3 20010522
                                                 WO 2001-KR845
                                                                            20010522
                                                 US 2002-958069
                                                                        A3 20020716
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ED Entered STN: 22 Nov 2002

AB Preparation and use of a catalyst for purification of diesel engine exhaust gas is presented, whereby the catalyst comprises a carrier of at least one sulfur-resistant refractory oxide and at least one catalytic metal, wherein at least one solid acid and/or H2SO4 is carried on the carrier by adding at least one precursor of said solid acid and/or H2SO4 during the preparation of the carrier, and preparation thereof. The refractory oxide is selected from: at least one oxide of Si, Al, Fe, Sn and/or Ce or their analogs in the form of a composite oxide or a mixture of oxides; zeolite; mordenite; and their mixts. The solid acid is selected from: tungsten oxides; molybdenum oxides; and their mixts.,. The catalytic metal is selected from: Pt, Pd, Rh, Ru, Re and their mixts.,. The catalyst preparation process comprises the steps of: (1) preparing a solution of at least one sulfur-resistant refractory oxide precursor; (2) adding an alkali solution such as an aqueous ammonia solution to co-precipitate and to form a gel or a mixed gel thereof; (3) drying,

shaping and calcining the resulted gel or mixed gel; and (4) depositing at least one catalytic metal. At least one solid acid precursor or its solution is added before or after the co-precipitation of step (2), and H2SO4 is added before or after the co-precipitation of step (2). The catalyst of this invention is thermally and chemical durable and can effectively remove the particulate matter, hydrocarbons and NOx contained in the diesel engine exhaust gas at low temps.

IT 7664-41-7, Ammonia, reactions

(for gelation of catalyst precursors; preparation and use of diesel engine exhaust catalyst including composite oxide and H2SO4)

CN · Ammonia (CA INDEX NAME)

NH3

IT 7440-06-4, Platinum, uses

(preparation and use of diesel engine exhaust catalyst including composite oxide and H2SO4)

RN 7440-06-4 HCAPLUS

CN Platinum (CA INDEX NAME)

Pt

IC ICM B01J021-06

ICS B01J029-89; B01J037-02

CC 59-3 (Air Pollution and Industrial Hygiene)
Section cross-reference(s): 67

ST diesel exhaust catalyst low temp composite oxide sulfuric acid; platinum tungsten zirconia titania composite oxide diesel exhaust catalyst

IT Refractory metal oxides

(sulfur- resistant; preparation and use of diesel engine exhaust catalyst including composite oxide and H2SO4)

IT 7664-41-7, Ammonia, reactions

(for gelation of catalyst precursors; preparation and use of diesel engine exhaust catalyst including composite oxide and H2SO4)

IT 1314-23-4D, Zirconia, composites with silica, titania, or tin oxide 7440-05-3, Palladium, uses 7440-06-4, Platinum, uses 7440-15-5, Rhenium, uses 7440-16-6, Rhodium, uses 7440-18-8, Ruthenium, uses 12028-48-7, Ammonium metatungstate 12173-98-7, Mordenite 13463-67-7D, Titania, composites with zirconia or tin oxide 18282-10-5D, Tin oxide sno2, composites with zirconia or titania

(preparation and use of diesel engine exhaust catalyst including composite oxide and H2SO4)

REFERENCE COUNT:

THERE ARE 5 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L68 ANSWER 8 OF 29 HCAPLUS COPYRIGHT 2007 ACS on STN

5

ACCESSION NUMBER:

2004:923812 HCAPLUS

DOCUMENT NUMBER:

142:80608

TITLE:

Treatment method of catalyst with water and gas for enhancing activity and reducing inactivation

INVENTOR(S):

Kim, Moon Chan; Son, In Hyuck

PATENT ASSIGNEE(S):

S. Korea

SOURCE:

Repub. Korean Kongkae Taeho Kongbo, No pp. given

CODEN: KRXXA7

DOCUMENT TYPE:

Patent Korean

LANGUAGE:
FAMILY ACC. NUM. COUNT:

1 /

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE	
KR 2002041346	 A	20020601	KR 2002-10338	20020226	
PRIORITY APPLN. INFO.:			KR 2002-10338	20020226	

Entered STN: 03 Nov 2004 ED

AB Provided is a refractory inorg. oxides catalyst supported by solid powder with enhanced activity and selectivity, exhibiting no inactivation after the use of 200 h. The catalyst is used in PROX reaction, elimination of volatile organic material and catalytic oxidation The treatment method comprises the steps of making inorg. organic catalyst such as alumina, titania and silica oxide by slurry washing to the honeycomb, drying, calcining at 300-800°C and cooling to 100°C; and improving the surface of the above catalyst by heating at more than 100°C and flowing on its surface with one or more liquid materials selected from water, alcs., ammonia water, hydrogen peroxide water and hydrochloride and with one or more gases chosen from hydrogen, oxygen, ozone, carbon monoxide, methane, propane and butane.

ICM B01J037-00 IC

67-1 (Catalysis, Reaction Kinetics, and Inorganic Reaction Mechanisms) CC Section cross-reference(s): 59

IT Oxides (inorganic), uses

Refractory metal oxides

(treatment method of catalyst with water and gas for enhancing activity and reducing inactivation)

L68 ANSWER 9 OF 29 HCAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER:

2002:370573 HCAPLUS

DOCUMENT NUMBER:

137:144487

TITLE:

Monitoring aging and deactivation of emission abatement catalysts for selective catalytic

reduction of NOx

AUTHOR(S):

Herman, Richard G.; Sale, John W.; Stenger, Harvey G., Jr.; Lyman, Charles E.; Agogliatti, John E.; Cai, Yeping; Ramachandran, Bala; Choi, Sukwon Zettlemoyer Center for Surface Studies, Lehigh

CORPORATE SOURCE:

University, Bethlehem, PA, 18015, USA

Topics in Catalysis (2002), 18(3-4), 251-257

SOURCE:

CODEN: TOCAFI; ISSN: 1022-5528

Kluwer Academic/Plenum Publishers

PUBLISHER: DOCUMENT TYPE: LANGUAGE:

Journal English

Entered STN: 19 May 2002

Titania/vanadia, zeolite, and noble metal catalysts are utilized for selective catalytic reduction (SCR) of NOx using ammonia as the reductant in different temperature ranges. Studies of aging have been carried out to probe deactivation rates and mechanisms. Periodic laboratory testing of samples of NOx reduction catalysts from multilayer reactors, such as those utilized at elec. power plants, allows prediction of catalyst lifetimes. Testing has been carried out under protocol conditions with monolith, plate-type, and pelleted catalysts so that relative NO reduction rates can be compared, with or without the presence of SO2. The catalysts were analyzed by surface anal. techniques, including electron microscopy and XPS, to probe surface morphol., loss of active components, presence of poisons, and blocking of pores and active sites by ammonium bisulfate to determine the dominant mode(s) of gradual deactivation.

IT 1314-62-1, Vanadia, uses 7440-06-4, Platinum, uses

(reduction catalysts containing; monitoring of aging and deactivation of catalysts for selective catalytic reduction of NOx by NH3)

RN 1314-62-1 HCAPLUS CN Vanadium oxide (V2O5) (CA INDEX NAME)

*** STRUCTURE DIAGRAM IS NOT AVAILABLE ***

RN 7440-06-4 HCAPLUS

CN Platinum (CA INDEX NAME)

Pt.

CC 59-4 (Air Pollution and Industrial Hygiene)

Section cross-reference(s): 51, 67

IT Mordenite-type zeolites

Zeolite ZSM-5

(Pt-containing, reduction catalysts; monitoring of aging and deactivation of catalysts for selective catalytic reduction of NOx by NH3)

IT 1314-62-1, Vanadia, uses 1344-28-1, Alumina, uses
7440-06-4, Platinum, uses 13463-67-7, Titania,
uses

(reduction catalysts containing; monitoring of aging and deactivation of catalysts for selective catalytic reduction of NOx by NH3)

REFERENCE COUNT:

24 THERE ARE 24 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE

RE FORMAT

L68 ANSWER 10 OF 29 HCAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER:

2001:885209 HCAPLUS

DOCUMENT NUMBER:

136:39097

TITLE:

Manufacture of heterogeneous catalysts

on micrometer range-particle supports
Roth, Marcel; Zander, Lars; Schwerin, Albrecht;

INVENTOR(S): Roth, Marcel; Zand

Gutsche, Bernhard

PATENT ASSIGNEE(S):

Henkel K.-G.a.A., Germany

SOURCE:

Ger. Offen., 8 pp.
CODEN: GWXXBX

DOCUMENT TYPE:

Patent

LANGUAGE:

German

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
DE 10025964	A1	20011206	DE 2000-10025964	20000525
PRIORITY APPLN. INFO.:			DE 2000-10025964	20000525

ED Entered STN: 07 Dec 2001

AB A title catalyst with increased surface comprises
Fe, Co, Ni, Cu, Ag, Au, Pd, Pt, Cd, Cr, Mn, W, V, Ti and/or
Mo preferably in oxide form, supported on particulate solid support
with particle size <100 μm. The support and, optionally, the
catalyst is magnetic or magnetizable. For example, a
dispersion of Fe oxide catalyst particles was prepared by
precipitation of FeCl3 and FeCl2·4H2O with aqueous ammonia containing
polyacrylic acid. The dispersion was dialyzed, concentrated by evaporation,

the

liquid concentrate combined with Novozym 435 and oleic acid-rich sunflower oil and the mixture treated at 60° with 70%-aqueous H2O2, the enzyme was separated by filtration and the catalyst separated by use of a magnetic field to give a product containing epoxidized oleic acid with 80%

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conversion. Nanoscale \gamma-Fe203 particles coated with
     WO3, V2O5 or MoO3 were also prepared
ΙT
     7440-06-4, Platinum, uses
        (manufacture of heterogeneous catalysts on micrometer-range
        magnetic particle supports)
     7440-06-4 HCAPLUS
RN
     Platinum (CA INDEX NAME)
CN
Pt.
     1314-62-1, Vanadium oxide (V2O5), uses
IT
        (particle shell; manufacture of heterogeneous catalysts on
       micrometer-range magnetic particle supports)
RN
     1314-62-1 HCAPLUS
     Vanadium oxide (V2O5)
CN
                           (CA INDEX NAME)
*** STRUCTURE DIAGRAM IS NOT AVAILABLE ***
IC
     ICM B01J023-00
     ICS B01J023-70; B01J031-26; C12N009-14; C07C409-24; C07C407-00;
          C07D301-12; C07C067-02; B01J023-85; B01J023-847
CC
     45-4 (Industrial Organic Chemicals, Leather, Fats, and Waxes)
ST
     catalyst epoxidn manuf magnetic particle support; iron oxide
     magnetic particle support epoxidn catalyst manuf; sunflower
     oil epoxidn iron oxide magnetic particle catalyst manuf
ΙT
     Alkenes, reactions
        (epoxidn.; manufacture of heterogeneous catalysts on
        micrometer-range magnetic particle supports)
IT
     Catalyst supports
       Epoxidation catalysts
       Oxidation catalysts
        (manufacture of heterogeneous catalysts on micrometer-range
        magnetic particle supports)
IT
     Carboxylic acids, preparation
        (peroxy, epoxidn. agents; manufacture of heterogeneous catalysts
        on micrometer-range magnetic particle supports)
IT
     Fatty acids, reactions
        (sunflower-oil, Me esters, epoxidn.; manufacture of heterogeneous
        catalysts on micrometer-range magnetic particle supports)
IT
     9001-62-1, Novozym 435
        (cocatalyst; manufacture of heterogeneous catalysts on
       micrometer-range magnetic particle supports)
IT
     7722-84-1, Hydrogen peroxide, uses
        (epoxidn. agent; manufacture of heterogeneous catalysts on
       micrometer-range magnetic particle supports)
ΙT
    111-66-0, 1-Octene
        (epoxidn.; manufacture of heterogeneous catalysts on
       micrometer-range magnetic particle supports)
IT
                           7439-96-5, Manganese, uses
     7439-89-6, Iron, uses
                                                         7439-98-7,
    Molybdenum, uses
                       7440-02-0, Nickel, uses
                                                  7440-05-3, Palladium,
    uses 7440-06-4, Platinum, uses
                                     7440-22-4,
                   7440-32-6, Titanium, uses
                                                7440-33-7, Wolfram, uses
    Silvèr, uses
     7440-43-9, Cadmium, uses
                               7440-47-3, Chromium, uses
                                                           7440-48-4,
                  7440-50-8, Copper, uses
     Cobalt, uses
                                             7440-57-5, Gold, uses
     7440-62-2, Vanadium, uses
        (manufacture of heterogeneous catalysts on micrometer-range
       magnetic particle supports)
     1332-37-2P, Iron oxide, preparation
        (manufacture of heterogeneous catalysts on micrometer-range
```

magnetic particle supports)

IT 1313-27-5, Molybdenum oxide (MoO3), uses 1314-35-8, Tungsten oxide (WO3), uses 1314-62-1, Vanadium oxide (V2O5), uses

13463-67-7, Titanium dioxide, uses

(particle shell; manufacture of heterogeneous catalysts on micrometer-range magnetic particle supports)

REFERENCE COUNT:

THERE ARE 5 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L68 ANSWER 11 OF 29 HCAPLUS COPYRIGHT 2007 ACS on STN

5

ACCESSION NUMBER:

2000:401547 HCAPLUS

DOCUMENT NUMBER:

133:26268

TITLE:

Resistance-based gas sensors with WO3-TiO2 active

layer for determination of NOx in automobile

exhaust gases

INVENTOR(S):

Kornely, Susanne; Seidl, Monika; Meixner, Hans;

Fleischer, Maximilian; Lampe, Uwe; Mrotzek,

Christine; Pohle, Roland; Giber, Janos Siemens Aktiengesellschaft, Germany

PATENT ASSIGNEE(S): SOURCE:

Eur. Pat. Appl., 10 pp.

CODEN: EPXXDW

Patent German

LANGUAGE:
FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

DOCUMENT TYPE:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE .
EP 1008847	A2	20000614	EP 1999-123914	19991201
EP 1008847	A3	20020605		
R: AT, BE, CH,	DE, DK	, ES, FR,	GB, GR, IT, LI, LU, NL	, SE, MC,
PT, IE, SI,	LT, LV	, FI, RO		
DE 19856369	A1	20000615	DE 1998-19856369	19981207
DE 19856369	C2	20001207		•
PRIORITY APPLN. INFO.:			DE 1998-19856369	A 19981207

ED Entered STN: 16 Jun 2000

AB A resistive gas sensor, especially suitable for detection of NO, NO2, NH3, or hydrocarbons in an automobile exhaust gas, consists of a gas-sensitive layer, a corresponding measuring electrode, and a heating unit, in which the gas-sensitive layer consists of a mixture of WO3 and TiO2, which is prepared by crystallizing WO3 around a nucleus of TiO2. The gas-sensitive layer (5-50 μm thick), which contains ≥50 weight% WO3, can be prepared by the sol-gel method using tungstic acid salt (M2WO4, in which M = H, Na, K, or NH4) precursors, or can be prepared from Ti(OC3H7)4 and WCl6 precursors. The gas sensor is also connected to an oxidation catalyst consisting of an impregnated metal oxide support (e.g., γ-Al2O3, SiO2, or TiO2 impregnated with a noble metal, such as Pt, Rh, Pd, or Ir) or a pure metal oxide catalyst (e.g., TiO2-V2O5 containing CuO or MnO2).

IT 7664-41-7, Ammonia, analysis

(determination of, in exhaust gases; resistance-based gas sensors with WO3-TiO2 active layer for determination of NOx in automobile exhaust gases)

RN 7664-41-7 HCAPLUS

CN Ammonia (CA INDEX NAME)

```
NH3
```

IT 1314-62-1, Vanadium oxide (V2O5), uses 7440-06-4,

Platinum, uses

(oxidation catalyst containing; resistance-based gas sensors with WO3-TiO2 active layer for determination of NOx in automobile exhaust gases)

RN 1314-62-1 HCAPLUS

CN Vanadium oxide (V2O5) (CA INDEX NAME)

*** STRUCTURE DIAGRAM IS NOT AVAILABLE ***

RN 7440-06-4. HCAPLUS

CN Platinum (CA INDEX NAME)

Pt

IC ICM G01N027-12

CC 79-2 (Inorganic Analytical Chemistry)
Section cross-reference(s): 59

IT Oxidation catalysts

(sensor containing; resistance-based gas sensors with WO3-TiO2 active layer for determination of NOx in automobile exhaust gases)

TT 7664-41-7, Ammonia, analysis 10102-43-9, Nitrogen oxide (NO), analysis 10102-44-0, Nitrogen oxide (NO2), analysis 11104-93-1, Nitrogen oxide, analysis

(determination of, in exhaust gases; resistance-based gas sensors with WO3-TiO2 active layer for determination of NOx in automobile exhaust gases)

IT 1313-13-9, Manganese oxide (MnO2), uses 1314-62-1, Vanadium oxide (V2O5), uses 1317-38-0, Copper oxide (CuO), uses 1344-28-1, Alumina, uses 7439-88-5, Iridium, uses 7440-05-3, Palladium, uses 7440-06-4, Platinum, uses 7440-16-6, Rhodium, uses 7631-86-9, Silica, uses

(oxidation catalyst containing; resistance-based gas sensors with WO3-TiO2 active layer for determination of NOx in automobile exhaust gases)

L68 ANSWER 12 OF 29 HCAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER:

2000:843199 HCAPLUS

DOCUMENT NUMBER:

134:90201

TITLE:

AUTHOR(S):

Multisensor system for remote detection of trace

gases in thin-layer metal oxide gas sensor arrays Wollenstein, J.; Jagle, M.; Scheulin, M.; Schmid,

J.; Bottner, H.; Becker, W. J.

CORPORATE SOURCE:

Freiburg, Germany

SOURCE:

VDI-Berichte (2000), 1530 (Sensoren und Messsysteme

2000), 191-200

CODEN: VDIBAP; ISSN: 0083-5560

PUBLISHËR:

VDI Verlag GmbH

DOCUMENT TYPE:

Journal German

LANGUAGE:

ED Entered STN: 03 Dec 2000

AB A gas sensor for remote measurements of trace gases was developed and tested. Thin (60-70 nm) layers of V2O5 and SnO2 were deposited on a 3+3 mm Si/SiO2 chip by vapor deposition and sputtering, resp., with subsequent annealing. The oxidic layers were optionally coated with a 1.5-nm layer of Pt as catalyst. The sensors were

tested by exposition to an 80:20 N/O mixture containing traces of CH4, NO2, CO, or NH3 (V20 with and without Pt catalyst) and CO, NO2, or O3 (SnO2 with or without Pt). In a long-term field test, the CO concentration was monitored in a road tunnel with a SnO2 sensor with Pt catalyst (to suppress cross-sensitivity to NO2).

IT 7664-41-7, Ammonia, analysis

(multisensor system for detection of trace gases in thinlayer metal oxide gas sensor arrays)

RN 7664-41-7 HCAPLUS

CN Ammonia (CA INDEX NAME)

NH3

IT 7440-06-4, Platinum, uses

(multisensor system for detection of trace gases in thin-layer metal oxide gas sensor arrays)

RN 7440-06-4 HCAPLUS

CN Platinum (CA INDEX NAME)

Pt

IT 1314-62-1, Vanadium oxide (V2O5), uses
 (multisensor system for detection of trace gases in thin-layer
 metal oxide gas sensor arrays)

RN 1314-62-1 HCAPLUS

CN Vanadium oxide (V2O5) (CA INDEX NAME)

*** STRUCTURE DIAGRAM IS NOT AVAILABLE ***

CC 59-1 (Air Pollution and Industrial Hygiene)
 Section cross-reference(s): 79

TT 74-82-8, Methane, analysis 630-08-0, Carbon monoxide, analysis 7664-41-7, Ammonia, analysis 10028-15-6, Ozone, analysis 10102-44-0, Nitrogen oxide (NO2), analysis (multisensor system for detection of trace gases in thinlayer metal oxide gas sensor arrays)

IT 7440-06-4, Platinum, uses

(multisensor system for detection of trace gases in thin-layer metal oxide gas sensor arrays)

IT 1314-62-1, Vanadium oxide (V2O5), uses 18282-10-5, Tin oxide (SnO2)

(multisensor system for detection of trace gases in thin-layer metal oxide gas sensor arrays)

REFERENCE COUNT:

THERE ARE 6 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L68 ANSWER 13 OF 29 HCAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER:

1999:142020 HCAPLUS

DOCUMENT NUMBER:

130:227135

TITLE: INVENTOR(S): Apparatus for biological treatment of garbage Mizobuchi, Manabu; Nakagawa, Shouji; Kinubawa,

Kensaku

PATENT ASSIGNEE(S):

Matsushita Electric Works, Ltd., Japan

SOURCE:

Jpn. Kokai Tokkyo Koho, 7 pp.

CODEN: JKXXAF

DOCUMENT TYPE:

Patent

LANGUAGE:

Japanese

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

PATENT NO.	· KIND	DATE	APPLICATION NO.	DATE
JP 11057669	Α	19990302	JP 1997-222645	19970819
PRIORITY APPLN. INFO.:			JP 1997-222645	19970819

ED Entered STN: 05 Mar 1999

AB The apparatus comprises (a) a biol. decomposition tank, (b) an air intake path, and (c) an exhaust path, and a purification apparatus comprising a metal chloride-adhered purification layer and a catalyst layer and a heater for the purification apparatus are placed at c. An ammonia adsorbing layer may be formed beneath the purification layer in the purification apparatus Odor generated during treatment of garbage is decreased.

IT 1314-62-1, Vanadium pentoxide, uses 7440-06-4,

Platinum, uses

(air purification and catalytic deodorization in apparatus for biol. treatment of garbage)

RN 1314-62-1 HCAPLUS

CN Vanadium oxide (V2O5) (CA INDEX NAME)

*** STRUCTURE DIAGRAM IS NOT AVAILABLE ***

RN 7440-06-4 HCAPLUS

CN Platinum (CA INDEX NAME)

Pt

IC ICM B09B003-00

ICS B01D053-86; B01J023-89

CC 60-4 (Waste Treatment and Disposal) Section cross-reference(s): 47, 59, 67

ST biol garbage treatment catalytic deodorization; metal chloride purifn layer garbage treatment; ammonia adsorption biol garbage treatment app

IT Zeolite-group minerals

(activated, ammonia adsorption layer; air purification and catalytic deodorization in apparatus for biol. treatment of garbage)

IT 1344-28-1, Alumina, uses

(activated, ammonia adsorption layer; air purification and catalytic deodorization in apparatus for biol. treatment of garbage)

IT 1313-99-1, Nickel oxide, uses 1314-62-1, Vanadium pentoxide, uses 1332-37-2, Iron oxide, uses 7440-05-3, Palladium, uses 7440-06-4, Platinum, uses 7440-16-6, Rhodium, uses 7440-18-8, Ruthenium, uses 7440-22-4, Silver, uses 7440-50-8, Copper, uses 7440-57-5, Gold, uses 11104-61-3, Cobalt oxide 11129-60-5, Manganese oxide

(air purification and catalytic deodorization in apparatus for biol. treatment of garbage)

L68 ANSWER 14 OF 29 HCAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER:

1997:191622 HCAPLUS

DOCUMENT NUMBER:

126:190273

TITLE:

Catalytic treatment of waste gases containing

harmful pollutants

INVENTOR(S):

Shimada, Takashi; Hatakeyama, Tosha; Nawa, Yoji

PATENT ASSIGNEE(S): Japan Pionics, Japan

SOURCE:

Jpn. Kokai Tokkyo Koho, 5 pp.

CODEN: JKXXAF

DOCUMENT TYPE:

Patent

1 .

LANGUAGE:

Japanese

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
	, - -			
JP 09000873	A·	19970107	JP 1995-174370	19950616
PRIORITY APPLN. INFO.:			JP 1995-174370	19950616

Entered STN: 22 Mar 1997 ED

Harmful pollutants (especially, amines or NH3) are removed from waste gases AB from semiconductor manufacturing by contacting with catalysts containing mainly CuO, MnO2, and Co(II) salts adhered on refractory metal oxide supports.

IT 7664-41-7, Ammonia, processes

(catalytic treatment of waste gases containing harmful pollutants)

RN7664-41-7 HCAPLUS

CN Ammonia (CA INDEX NAME)

NH₃

IC ICM B01D053-58

ICS B01D053-72; B01J020-06; B01J023-889

CC 59-4 (Air Pollution and Industrial Hygiene)

74-89-5, Monomethylamine, processes 75-50-3, Trimethylamine, processes 124-40-3, Dimethylamine, processes 302-01-2, Hydrazine, processes 7664-41-7, Ammonia, processes 30260-66-3, Dimethylhydrazine

(catalytic treatment of waste gases containing harmful pollutants)

L68 ANSWER 15 OF 29 HCAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER:

1997:307657 HCAPLUS

DOCUMENT NUMBER:

126:282028

TITLE:

Layered catalysts for exhaust gas treatment

INVENTOR(S): PATENT ASSIGNEE(S): Morsbach, Bernd

BASF A.-G., Germany

SOURCE:

Eur. Pat. Appl., 7 pp. CODEN: EPXXDW

DOCUMENT TYPE:

Patent

LANGUAGE:

German

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
EP 763380	A1	19970319	EP 1996-114446	19960910
R: BE, DE, FR,	GB, NL		•	
DE 19534497	A1	19970320	DE 1995-19534497	19950918
PRIORITY APPLN. INFO.:			DE 1995-19534497 A	19950918

ED Entered STN: 14 May 1997

AB The layered catalysts comprise ≥1 inner and ≥1

outer layer, where the center or the inner layers comprise oxide layers containing noble metals (e.g., Pt , Pd, and/or Rh) and the outer layers contain the components A and C or A, B, and C, where A is an oxide of the elements Ti, Al, Zr, or their mixts., B is an oxide of the elements Mo, W, or their mixts., and C is an oxide or sulfate of the elements V, Fe, Mn; Ni, Co, Cu, Nb, Zn, or their mixts. and where the carrier center and each layer may optionally also contain an oxide or sulfate of the elements Si, B, Zn, or their mixts., inorg. fibers, clays or their mixts. The carrier may be ≥1 component chosen from A, B, or C, or their mixts., and inert carrier (e.g., cordierite or mullite) or a metallic carrier with and least one oxidic support layer. The catalyst is especially suitable for removal of nitrogen oxides, carbon monoxide, and hydrocarbons from oxygen-containing exhaust gas at 50-800° and 0.01-200 bar using ammonia or and ammonia releasing agent as reductant. 1314-62-1, Vanadium oxide, uses 7440-06-4, Platinum, uses (layered catalysts for exhaust gas treatment) 1314-62-1 HCAPLUS Vanadium oxide (V2O5) (CA INDEX NAME) * * * STRUCTURE DIAGRAM IS NOT AVAILABLE *** 7440-06-4 HCAPLUS Platinum (CA INDEX NAME) 7664-41-7, Ammonia, reactions (reductant; layered catalysts for exhaust gas treatment) 7664-41-7 HCAPLUS Ammonia (CA INDEX NAME) ICM B01J035-00 ICS B01J037-02; B01D053-86; B01D053-94 59-3 (Air Pollution and Industrial Hygiene) Section cross-reference(s): 51, 67 1313-27-5, Molybdenum oxide, uses 1313-96-8, Niobium oxide 1313-99-1, Nickel oxide, uses 1314-13-2, Zinc oxide; uses 1314-23-4, Zirconium oxide, uses 1314-35-8, Tungsten oxide, uses 1314-62-1, Vanadium oxide, uses 1332-37-2, Iron oxide, uses 1344-28-1, Aluminum oxide (Al2O3), uses 1344-70-3, Copper oxide 7440-05-3, Palladium, uses 7440-06-4, Platinum, 7440-16-6, Rhodium, uses 7733-02-0, Zinc sulfate 7785-87-7, Manganese sulfate 7786-81-4, Copper sulfate, uses Nickel sulfate 10124-43-3, Cobalt sulfate 10124-49-9, Iron sulfate 11104-61-3, Cobalt oxide 11129-60-5, Manganese oxide 13463-67-7, 16785-81-2, Vanadium sulfate Titanium oxide, uses 36220-20-9, Niobium sulfate (layered catalysts for exhaust gas treatment) 7664-41-7, Ammonia, reactions (reductant; layered catalysts for exhaust gas treatment)

IT

RN

CN

RN

CN

Pt

IT

RN

CN

NH₃

IC

CC

IT

IT

ANSWER 16 OF 29 HCAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER:

1996:664605 HCAPLUS

DOCUMENT NUMBER:

125:281737

TITLE:

Porous sintered steel infiltrated with low-density metals for sliding parts resistant to seizing Fujine, Manabu; Kajikawa, Yoshiaki; Yamashita,

INVENTOR(S):

Minoru; Saito, Koji

PATENT ASSIGNEE(S):

Toyota Jidosha Kabushiki Kaisha, Japan

SOURCE:

Eur. Pat. Appl., 27 pp.

CODEN: EPXXDW Patent

DOCUMENT TYPE: LANGUAGE:

English

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
EP 732417	A1	19960918	EP 1996-104256	19960318
EP 732417	B1	20020213		
R: DE, FR, GB,	IT, SE		•	•
JP 08319504	A	19961203	JP 1996-56518	. 19960313
JP 3191665	B2	20010723		
AU 9648135	A	19960926	AU 1996-48135	19960315
AU 710033	· B2	19990909·		
KR 183227	B1	19990401	KR 1996-6946	19960315
CA 2172029	С	20010515	CA 1996-2172029	19960318
PRIORITY APPLN. INFO.:			JP 1995-59455	A 19950317
		•	JP 1996-56518	19960313

ED Entered STN: 11 Nov 1996

The composites for sliding parts are manufactured from sintered porous steel (or Fe alloy) having Vickers microhardness of 200-800, and are infiltrated with low-d. metal (especially Al or Mg alloys) for increased resistance to seizing. The sintered steels or Fe alloys have nominal d. at 30-88% of theor., and optionally contain dispersed hard particles (especially carbides) at ≤50 volume%. The sintered alloy steels typically contain C 0.5-1.2, Cr 5.8-8.7, Mo 0.1-0.6, and V 0.1-0.6 weight%. The Al alloy for infiltration is typically molten AC8A alloy nominally containing Cu 0.8-1.3, Si 11-13, and Mg 0.7-1.3 weight%, and can be heat treated for age hardening after the infiltration of sintered parts. The sintered steel having 60% of theor. d. was manufactured from the atomized Fe-0.2 C-1 Si-0.4 Mn-5 Cr-1.3 Mo-1 weight% V steel powder of SKD61 type, infiltrated with Al-alloy melt, and showed no seizing in a sliding test against nitrided steel at 250°.

IT 7664-41-7D, Ammonia, dissociated

> (cooling in, of sintered parts; sintered steel parts cooled in gas and infiltrated with low-d. metal for resistance to seizing)

RN 7664-41-7 HCAPLUS

Ammonia (CA INDEX NAME) CN

NH₃

IT

(hard phase, dispersed; sintered steel hardened with dispersed particles and infiltrated with low-d. metal for resistance to seizing)

RN11122-73-9 HCAPLUS

CN Chromium alloy, nonbase, Cr, Fe (CA INDEX NAME) Component Component
Registry Number
Cr 7440-47-3
Fe 7439-89-6

IC ICM C22C033-02

CC 55-4 (Ferrous Metals and Alloys)

IT Aluminum alloy, base Magnesium alloy, base

(infiltration with molten; sintered steel parts infiltrated with low-d. alloys for resistance to seizing)

IT Iron alloy, base

(sintered, sliding parts; porous steel infiltrated with low-d. metal for sliding parts resistant to seizing)

IT 1333-74-0, Hydrogen, processes **7664-41-7D**, Ammonia , dissociated 7727-37-9, Nitrogen, processes

(cooling in, of sintered parts; sintered steel parts cooled in gas and infiltrated with low-d. metal for resistance to seizing)

IT 11122-73-9 12783-13-0 60719-59-7, Chromium iron carbide (hard phase, dispersed; sintered steel hardened with dispersed particles and infiltrated with low-d. metal for resistance to seizing)

L68 ANSWER 17 OF 29 HCAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER:

1996:153530 HCAPLUS

DOCUMENT NUMBER:

124:184317

TITLE:

Method for denitrating exhaust gases

INVENTOR (S):

Iida, Kouzo; Nojima, Shigeru; Obayashi, Yoshiaki;

Kobayashi, Norihisa; Serizawa, Satoru

PATENT ASSIGNEE(S):

Mitsubishi Jukogyo Kabushiki Kaisha, Japan Eur. Pat. Appl., 14 pp.

SOURCE:

CODEN: EPXXDW

DOCUMENT TYPE:

Patent

LANGUAGE:

English

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.		DATE
EP 694329	A2	19960131	EP 1995-111683	•	19950725
EP 694329	A3	19970813			
EP 694329	B1	20001018			
R: AT, DE, IT,	NL		•		
JP 08038856	Α	19960213	JP 1994-176494		19940728
JP 3462580	. B2	20031105			
JP 08103633	Α	19960423	JP 1994-238892		19941003
JP 3241216	B2	20011225			
CA 2154500	A1	19960129	CA 1995-2154500		19950724
CA 2154500	С	20011002			
AT 196998	T	20001115	AT 1995-111683		19950725
US 5728356	Α	19980317	US 1995-508174		19950727
US 6080376	Α	20000627	US 1997-988116		19971210
PRIORITY APPLN. INFO.:			JP 1994-176494	Α	19940728
•					
			JP 1994-238892	Α	19941003
			US 1995-508174	А3	19950727

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Entered STN: 16 Mar 1996
ED
     Nitrogen oxides are catalytically removed using ammonia as a
AB
     reducing agent in the presence of a catalyst comprising a denitration
     catalyst layer in the upstream of the gas flow, an
     ammonia decomposition catalyst layer capable of decomposing
     ammonia into nitrogen oxides in the downstream and a 2nd
     denitration catalyst layer or a denitration catalyst
     layer capable of decomposing ammonia in the further
     downstream. Ammonia is used at an amount of not less than the
     reaction equivalent for the nitrogen oxides in the exhaust gas .
     1314-62-1, Vanadium pentoxide, uses 7440-06-4,
IT
     Platinum, uses
        (method for denitrating exhaust gases).
     1314-62-1 HCAPLUS
RN
                            (CA INDEX NAME)
CN
     Vanadium oxide (V2O5)
***
    STRUCTURE DIAGRAM IS NOT AVAILABLE ***
RN
     7440-06-4 HCAPLUS
CN
     Platinum (CA INDEX NAME)
Pt
IC
     ICM B01D053-86
     ICS B01J029-04
CC
     59-3 (Air Pollution and Industrial Hygiene)
     1314-35-8, Tungsten trioxide, uses 1314-62-1, Vanadium
IT
     pentoxide, uses 7439-88-5, Iridium, uses
                                                 7439-91-0, Lanthanum,
           7440-03-1, Niobium, uses 7440-05-3, Palladium, uses
                               7440-16-6, Rhodium, uses
     7440-06-4, Platinum, uses
                                7440-32-6, Titanium, uses
     7440-18-8, Ruthenium, uses
                                                              7440-36-0,
                    7440-45-1, Cerium, uses 7440-47-3, Chromium, uses
     Antimony, uses
                                                         7440-62-2,
                              7440-55-3, Gallium, uses
     7440-48-4, Cobalt, uses
     Vanadium, uses 13463-67-7, Titania, uses
        (method for denitrating exhaust gases)
L68 ANSWER 18 OF 29 HCAPLUS COPYRIGHT 2007 ACS on STN
ACCESSION NUMBER:
                         1996:80488 HCAPLUS
DOCUMENT NUMBER:
                         124:124719
                         Chemical vapor deposition of silicon nitride
TITLE:
                         filaments from silicon subhydrides and
                         ammonia
AUTHOR (S):
                         Linner, Britta; Guggenberger, Michael A.;
                         Huettinger, Klaus J.; Kleebe, Hans-Joachim
CORPORATE SOURCE:
                         Inst. Chem. Tech., Univ. Karlsruhe, Karlsruhe,
                         D-76128, Germany
SOURCE:
                         Journal of the European Ceramic Society (1996),
                         16(1), 15-23
                         CODEN: JECSER; ISSN: 0955-2219
PUBLISHER:
                         Elsevier
DOCUMENT TYPE:
                         Journal
LANGUAGE:
                         English '
    Entered STN: 07 Feb 1996
AB
     This paper describes the synthesis of monocryst. \alpha-silicon
    nitride filaments. The synthesis is based on a catalyzed
     chemical vapor deposition process using iron or iron alloys as catalysts
     and silicon subhydrides and ammonia as gaseous precursors of
     silicon nitride. For in situ production of silicon subhydrides by
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gasification of silicon powder with hydrogen superficially nitrided

silicon powder was used to guarantee constant production rates up to 10 h and more. The kinetics of filament growth are shown to be determined by the solubility of nitrogen in and the diffusion of nitrogen through the catalyst particle.

IT 11122-73-9

(catalysts; CVD of silicon nitride filaments from silicon subhydrides and ammonia using iron or iron alloys as catalysts)

RN 11122-73-9 HCAPLUS

CN Chromium alloy, nonbase, Cr,Fe (CA INDEX NAME)

Component Component Registry Number

Cr 7440-47-3 Fe 7439-89-6

IT 7664-41-7, Ammonia, processes

(precursor; CVD of silicon nitride filaments from silicon subhydrides and ammonia using iron or iron alloys as catalysts)

RN 7664-41-7 HCAPLUS

CN Ammonia (CA INDEX NAME)

NH3

CC 57-2 (Ceramics)

IT Vapor deposition processes

(CVD of silicon nitride filaments from silicon subhydrides and ammonia using iron or iron alloys as catalysts)

IT Crystal whiskers

(silicon nitride; CVD of silicon nitride filaments from silicon subhydrides and ammonia using iron or iron alloys as catalysts)

TT 7439-89-6, Iron, uses 11110-23-9 11122-73-9
(catalysts; CVD of silicon nitride filaments from silicon subhydrides and ammonia using iron or iron alloys as catalysts)

IT 7664-41-7, Ammonia, processes 50808-20-3, Silicon
hydride

(precursor; CVD of silicon nitride filaments from silicon subhydrides and ammonia using iron or iron alloys as catalysts)

IT 12033-89-5P, Silicon nitride, preparation
 (whiskers; CVD of silicon nitride filaments from silicon
 subhydrides and ammonia using iron or iron alloys as
 catalysts)

L68 ANSWER 19 OF 29 HCAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER:

1995:989089 HCAPLUS

DOCUMENT NUMBER:

124:69673

TITLE:

Voltammetry in the absence of a solution phase with solids prepared by a sol-gel process as the electrolytes: facilitation of an electrocatalytic anodic process in the presence of ammonia

anodic process in the presence of ammonia

AUTHOR(S): Cox

Cox, James A.; Alber, Kathryn S.; Tess, Mark E.;

Cummings, T. E.; Gorski, Waldemar

CORPORATE SOURCE: Department of Chemistry, Miami University, Oxford,

OH, 45056, USA

SOURCE:

LANGUAGE:

Journal of Electroanalytical Chemistry (1995),

396(1-2), 485-90

CODEN: JECHES

PUBLISHER:
DOCUMENT TYPE:

Elsevier Journal English

ED Entered STN: 19 Dec 1995

An interdigitated microelectrode (IME) coated with a glassy polymer of V2O5 by a sol-gel process is demonstrated to serve as a solid electrolyte for voltammetric studies in the absence of a contacting solution phase. The oxidation of Fe(II)-1,10-phenanthroline immobilized therein occurs at the same potential as in solution-phase expts. at a Pt working electrode; however, the current limiting process in the solid-state system depends on the time scale of the experiment Cyclic voltammetry at scan rates of 0.1-1.0 V/s yields currents limited by planar diffusion; but at <3 mV/s the peak currents are independent of scan rate. This steady-state behavior in the coated IME is indicative of current limitation by semi-cylindrical diffusion to the 10 µm + 5 mm Pt surfaces at slow scan rates; potential-step chronoamperometry verifies this interpretation. When 1 set of Pt fingers in the IME is a quasi-reference and the other set is modified with a polymeric Ru oxide catalyst, the presence of NH3 in the surrounding gas phase causes an anodic process. Indirect evidence that this process is the electrocatalytic oxidation of NH3 is presented. This anodic behavior is not observed when the voltammetry is performed in a conventional solution cell under otherwise-identical conditions.

IT 1314-62-1, Vanadium pentoxide, uses

(glassy polymer; interdigitated microelectrode coated with glassy polymer of vanadium pentoxide and facilitation of electrocatalytic anodic process in presence of ammonia)

RN 1314-62-1 HCAPLUS

CN Vanadium oxide (V2O5) (CA INDEX NAME)

*** STRUCTURE DIAGRAM IS NOT AVAILABLE ***

IT 7440-06-4, Platinum, uses

(voltammetry in absence of solution phase with solids prepared by sol-gel process as electrolytes and facilitation of electrocatalytic anodic process at ruthenia-modified platinum in presence of ammonia)

RN 7440-06-4 HCAPLUS

CN Platinum (CA INDEX NAME)

Pt

CC 72-2 (Electrochemistry)

Section cross-reference(s): 67

IT Oxidation catalysts

(electrochem., voltammetry in absence of solution phase with solids prepared by sol-gel process as electrolytes and facilitation of electrocatalytic anodic process at ruthenia-modified platinum in presence of ammonia)

IT Polyoxyalkylenes, uses

(fluorine- and sulfo-containing, ionomers, interdigitated microelectrode coated with Nafion and facilitation of electrocatalytic anodic process in presence of ammonia)

IT Fluoropolymers

(polyoxyalkylene-, sulfo-containing, ionomers, interdigitated microelectrode coated with Nafion and facilitation of electrocatalytic anodic process in presence of ammonia)

IT Ionomers

(polyoxyalkylenes, fluorine- and sulfo-containing, interdigitated microelectrode coated with Nafion and facilitation of electrocatalytic anodic process in presence of ammonia)

IT 1314-62-1, Vanadium pentoxide, uses

(glassy polymer; interdigitated microelectrode **coated** with glassy polymer of vanadium pentoxide and facilitation of electrocatalytic anodic process in presence of **ammonia**)

IT 11113-84-1, Ruthenium oxide

(mixed-valence cyano cross-linked polymeric; voltammetry in absence of solution phase with solids prepared by sol-gel process as electrolytes and facilitation of electrocatalytic anodic process at ruthenia-modified platinum in presence of ammonia)

IT 7440-06-4, Platinum, uses

(voltammetry in absence of solution phase with solids prepared by sol-gel process as electrolytes and facilitation of electrocatalytic anodic process at ruthenia-modified platinum in presence of ammonia)

L68 ANSWER 20 OF 29 HCAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER:

1991:589024 HCAPLUS

DOCUMENT NUMBER:

115:189024

TITLE:

Apparatus for treatment of diesel exhaust gases

INVENTOR(S):

Kawamura, Satoshi

PATENT ASSIGNEE(S):

Mitsubishi Heavy Industries, Ltd., Japan

SOURCE:

Jpn. Kokai Tokkyo Koho, 4 pp. CODEN: JKXXAF

DOCUMENT TYPE:

Patent

LANGUAGE:

Japanese

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
JP 03130522	. A	19910604	JP 1989-263994	19891012
PRIORITY APPLN. INFO.:			JP 1989-263994	19891012

ED Entered STN: 01 Nov 1991

AB NOx is removed from diesel exhaust gases by catalytic reduction with NH3 in an apparatus comprising means for injecting NH3 into the down stream of diesel engine exhaust duct, means for passing the gas mixture through a porous ceramic filter loaded with catalysts (e.g., V2O5-TiO2) for decomposing NOx into N2 and H2O, means for backwashing the filter and catalytically combusting the trapped dust and tar, and means for controlling the temperature of catalyst bed and preventing the pressure loss in the ceramic filter.

IT 1314-62-1, Vanadium oxide (V2O5), uses and miscellaneous 7440-06-4, Platinum, uses and miscellaneous

(catalyst containing, on porous ceramic filter, for diesel exhaust gas treatment)

RN 1314-62-1 HCAPLUS

CN Vanadium oxide (V2O5) (CA INDEX NAME)

*** STRUCTURE DIAGRAM IS NOT AVAILABLE ***

RN 7440-06-4 HCAPLUS

CN Platinum (CA INDEX NAME)

Pt

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IC ICM F01N003-08
ICS B01D053-36
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CC 59-4 (Air Pollution and Industrial Hygiene)

IT 1314-35-8, Tungsten oxide, uses and miscellaneous 1314-62-1,
 Vanadium oxide (V2O5), uses and miscellaneous 7440-05-3, Palladium,
 uses and miscellaneous 7440-06-4, Platinum, uses
 and miscellaneous 11098-99-0, Molybdenum oxide
 (catalyst containing, on porous ceramic filter, for diesel exhaust

(catalyst containing, on porous ceramic filter, for diesel exhaust gas treatment)

IT 11104-93-1, Nitrogen oxide, uses and miscellaneous (removal of, from diesel exhaust gases, by catalytic reduction with ammonia, on catalyst-coated porous ceramic filter)

L68 ANSWER 21 OF 29 HCAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER:

1986:55593 HCAPLUS

DOCUMENT NUMBER:

104:55593

TITLE:

Vanadium oxide catalyst for nitrogen oxide

reduction and its use in a process

INVENTOR(S):

Heck, Ronald M.; Keith, Carl D.; Farrauto, Robert

J.

PATENT ASSIGNEE(S):

SOURCE:

Engelhard Corp., USA

Eur. Pat. Appl., 17 pp.
CODEN: EPXXDW

DOCUMENT TYPE:

LANGUAGE:

Patent

English

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

PA	TENT NO.	KIND	DATE	APPLICATION NO.		DATE
			÷		-	
EP	161743	A2	19851121	EP 1985-301150		19850221
EP	161743	A3	19860402	•		
EP	161743	B1	19880928			•
	R: AT, BE, CH,	, DE, FR	R, GB, IT,	LI, LU, NL, SE		•
CA	1238628	A1	19880628	CA 1985-474859		19850221
AT	37490	T	19881015	AT 1985-301150		19850221
PRIORITY	Y APPLN. INFO.;			US 1984-582368	A	19840222
				EP 1985-301150	A	19850221

ED Entered STN: 23 Feb 1986

AB A catalyst for selective catalytic reduction of NOx with NH3 in a waste gas stream, preventing P-contamination of the catalyst, has an upstream section comprising a P-retaining material and a downstream section comprising a catalyst containing an effective amount of V2O5, e.g., 0.5-15 weight% on a refractory metal oxide support, e.g. Al2O3 or TiO2. The P-retaining material, e.g. activated may also be on a refractory support.

IT 7664-41-7, uses and miscellaneous

(nitrogen oxide reduction with, in waste gases, catalyst phosphorus contamination prevention in)

RN 7664-41-7 HCAPLUS

CN Ammonia (CA INDEX NAME)

NH3

IC ICM B01D053-36

CC 59-4 (Air Pollution and Industrial Hygiene)

Section cross-reference(s): 49, 51, 67

ST vanadium oxide composite redn catalyst; phosphorus contamination prevention redn catalyst; nitrogen oxide catalytic redn ammonia

IT 7664-41-7, uses and miscellaneous

(nitrogen oxide reduction with, in waste gases, catalyst phosphorus contamination prevention in)

IT 11104-93-1, uses and miscellaneous

(removal of, from waste gases, catalytic reduction with ammonia for, catalyst phosphorus contamination prevention in)

L68 ANSWER 22 OF 29 HCAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER:

1982:14262 HCAPLUS

DOCUMENT NUMBER:

96:14262

TITLE:

Platinum thin film resistance element

INVENTOR(S): Ohno, Yoshio

PATENT ASSIGNEE(S):

Kirk K. K., Japan

SOURCE:

Eur. Pat. Appl., 42 pp.

CODEN: EPXXDW

DOCUMENT TYPE:

Patent

LANGUAGE:

English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

	PATENT NO.	KIND	DATE	APPLICATION NO.	1	DATE
	EP 38078	A1	19811021	EP 1981-102856	-	19810414
	EP 38078	B1	19850313			•
	R: DE, FR, GB,	NL				•
	JP 56147048	Ą	19811114	JP 1980-49205		19800416
	JP 56147049	Α	19811114	JP 1980-49206		19800416
	JP 56147050	Α	19811114	JP 1980-49208		19800416
	JP 56150339	Α	19811120	JP 1980-49207		19800416
	JP 57101750	Α	19820624	JP 1980-177220		19801217
	EP 63264	A1	19821027	EP 1982-102641	•	19810414
	EP 63264	B1	19841212			
	R: DE, FR, GB,	NL				
	GB 2110165	Α	19830615	GB 1981-36331		19811202
	GB 2110165	В	19850911			•
PRIC	RITY APPLN. INFO.:			JP 1980-49204	Α	19800416
	·*					
				JP 1980-49205	Α	19800416
	•			•		
		r		JP 1980-49206	Α	19800416
			•	•		
				JP 1980-49207	Α	19800416
				•		
			•	JP 1980-49208	Α	19800416
				•		
				JP 1980-177220	Α	19801217
			•	EP 1981-102856	Α	19810414

ED Entered STN: 12 May 1984

AB A method is described for preparing a stable Pt thin-film

high-resistance resistor which does not require a 3 or 4 core lead wire and which can be used as an accurate temperature sensor and a gas sensor for low concns. of CO, NO, NH3, or an inflammable gas. The Pt film resistor is formed by sputtering on an insulator substrate which is stable at ≤1000°, preferably a cylinder or column, a 200-1000 Å Pt film at a power of 0.8 W/cm2, heat aging by raising the temperature in steps of 100° to 1000°, forming a spiral kerf in the film to obtain the desired resistance, and attaching lead wires to both ends of the film. forming a temperature sensor, the film is covered with an insulating polyimide or silicone film. In forming a gas sensor for CO, a thin layer of Cu oxide is deposited on the Pt, for NO detection a thin layer of a rare earth oxide 10-30, AgNO3 0.5-5 weight %, and balance V2O5 is deposited, for NH3 detection a layer of rare earth oxide 3-10, Sb203 1-5, AgN03 0.5-5 weight %, and balance V205, and for sp. flammable gas detection an Al2O3 or BeO cement is interposed between the catalytic metal oxide semiconductor and the Pt. 1314-62-1, uses and miscellaneous (catalyst, in oxide coating on platinum resistor for gas sensors) 1314-62-1 HCAPLUS Vanadium oxide (V2O5) (CA INDEX NAME) STRUCTURE DIAGRAM IS NOT AVAILABLE *** 7664-41-7, analysis (detection of, rare earth oxide-antimony oxide-silver nitrate-vanadium pentoxide coated platinum resistor sensor for) 7664-41-7 HCAPLUS Ammonia (CA INDEX NAME) 7440-06-4, uses and miscellaneous (resistor, thin-film, sputter-deposition of) 7440-06-4 HCAPLUS Platinum (CA INDEX NAME) CN · H01C007-22; H01C017-12; H01C013-00; G01N027-12; G01K007-18 76-2 (Electric Phenomena) Section cross-reference(s): 67, 75, 79, 80 sputtering platinum film resistor; temp sensor platinum resistor film; gas sensor platinum resistor film; catalytic semiconductor oxide gas sensor Gas analysis

IT

RN

CN

IT

RN

CN

NH3

IT

RN

Pt

IC

CC ST IT (detection of, semiconductor catalytic oxideplatinum film resistor sensor for) IT Rare earth oxides (gas sensors from sputtered platinum resistors coated with) IT Sputtering (of platinum thin-film resistor) TT Catalysts and Catalysis

(semiconductor oxide, for gas detection, on thin-film

platinum resistor support) IT Temperature (sensors for, from sputtered platinum film coated with silicone or polyimide) IT Polyimides, uses and miscellaneous Siloxanes and Silicones, uses and miscellaneous (temperature sensors from platinum resistor coated with) IT Electric resistors (film, platinum, sputter-deposition of) IT Combustibles (gaseous, detection of, semiconductor oxide-cement coated platinum resistor sensor for) IT 1309-64-4, uses and miscellaneous (catalyst, in oxide coating for platinum resistor for ammonia gas sensor) IT 7761-88-8, uses and miscellaneous (catalyst, in oxide coating on platinum film resistor for gas sensors) 1314-62-1, uses and miscellaneous IT (catalyst, in oxide coating on platinum resistor for gas sensors) IT 1344-70-3 (catalyst, on platinum resistor for carbon monoxide gas sensor) 630-08-0, analysis TT (detection of, copper oxide-coated sputtered platinum resistor sensor for) IT 7664-41-7, analysis (detection of, rare earth oxide-antimony oxide-silver nitrate-vanadium pentoxide coated platinum resistor sensor for) 10102-43-9, analysis IT (detection of, rare earth oxide-silver nitrate-vanadium pentoxide coated platinum resistor sensor for) IT 1344-28-1, uses and miscellaneous 1304-56-9 (flammable gas sensor from oxide coated platinum resistor with intermediate layer of) IT7440-06-4, uses and miscellaneous (resistor, thin-film, sputter-deposition of) L68 ANSWER 23 OF 29 HCAPLUS COPYRIGHT 2007 ACS on STN 1969:89301 HCAPLUS ACCESSION NUMBER: DOCUMENT NUMBER: 70:89301 Efficient and economical catalytic TITLE: oxidation of ammonia in the production of nitric oxide Keith, Carl D. INVENTOR(S): PATENT ASSIGNEE(S): Engelhard Minerals and Chemicals Corp. SOURCE: U.S., 7 pp. CODEN: USXXAM DOCUMENT TYPE: Patent LANGUAGE: English FAMILY ACC. NUM. COUNT: PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.		DATE
				-	
US 3428424	A	19690218	US 1965-434759		19650224
PRIORITY APPLN. INFO.:			US 1965-434759	Α	19650224

```
Entered STN: 12 May 1984
ED
     NO is prepared by reacting gaseous NH3 and atmospheric air in the presence of a
AB
     catalyst which is prepared by depositing 1-10% of a
     catalytic metal, such as Pt, Rh, Ir or alloys of
     Pt with Rh, Pd, or Ir, onto the gas flow channels of a porous
     inert unitary refractory skeletal structure previously coated with a
     catalytically active refractory metal
     oxide. The NO is then oxidized to NO2 in a nitric acid plant
     and the NO2 absorbed in H2O to form HNO3. The catalyst is
     kept at 650-1000° and 14-110 psig. The skeletal structure is
     prepared, e.g., from zirconmullite, and the refractory
     metal oxide prepared, e.g., by calcining hydrous
     alumina at 300-800°, is deposited as a continuous thin film of
     0.0004-0.001-in. thickness. Thus, such a catalyst containing 2%
    of Pt group metal consisting of an alloy of Rh 20 and
     Pt 80%, dispersed on the surfaces of gas flow
     channels and superficial macropores in contact with a corrugated
     porous refractory ceramic cylinder of \alpha-Al203, was placed in an
     NH3 converter. The corrugated porous cylinder had a diameter of 3.875
     in., was 1.875-in. long with 10 corrugations per in. which defined 20
     straight-through unobstructed gas flow channels per in. NO was prepared
     by passing a mixture of gaseous anhydrous NH3 1 and air 9 parts by volume,
     preheated to 200°, through the converter where the
     catalyst was at 925° and 110 psig. The
     catalyst showed excellent activity for oxidizing NH3 to NO.
     In comparison with the conventional process, 1/10 of the
     catalytic metal content can be employed and a materially lower
     pressure drop is obtained with this new invention. Also, the
     catalyst of this process enabled a weight hourly space velocity
     of NH3 of .apprx.10 times that of the conventional catalyst.
     C01B021-26A
TC
INCL 023162000
     49 (Industrial Inorganic Chemicals)
CC
     ammonia oxidn; oxidn ammonia; nitric acid prodn;
ST
     catalyst nitric acid prodn; platinum nitric acid
     prodn; rhodium nitric acid prodn
IT
     Rhodium alloys, containing
        (platinum-, as oxidation catalysts for
        ammonia)
IT
     Oxidation catalysts
        (platinum-rhodium alloy skeletal, for ammonia)
IT
     Platinum alloys, base
        (rhodium-, as oxidation catalysts for ammonia)
     10102-43-9P, preparation
IT
        (from ammonia, platinum-rhodium alloy oxidation
        catalysts for)
=> d 24-25 full
                                                THE THOMSON CORP on STN
    ANSWER 24 OF 29 WPIX COPYRIGHT 2007
L68
     1994-357967 [44]
AN
                        WPIX
    C1994-163322 [44]
DNC
     Removing metal carbonyls from a gas stream especially synthesis gas - by
     contact with lead oxide on a support
DC
     E17; E37; H04; J01
IN
     CARR N L
     (DENO-C) DEN NORSKE STATS OLJESELSKAP AS; (DENO-C) STATOIL DEN NORSKE
PA
     STATS OLJESELSKAP AS
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CYC 46

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PΙ
     WO 9425142
                     Al 19941110 (199444) * EN
                                               21.[0]
                     A · 19941121 (199508)
                                           EN
     AU 9466916
     US 5451384
                     Α
                        19950919 (199543)
                                           EN
                                               6[0]
     NO 9504203
                     Α
                        19951020 (199602)
                                           NO
     NO 180570
                     В
                       19970203 (199712)
                                           NO
     WO 9425142 A1 WO 1994-NO78 19940421; US 5451384 A US 1993-52395
ADT
     19930423; AU 9466916 A AU 1994-66916 19940421; NO 9504203 A WO
     1994-NO78 19940421; NO 180570 B WO 1994-NO78 19940421; NO 9504203 A NO
     1995-4203 19951020; NO 180570 B NO 1995-4203 19951020
FDT NO 180570 B Previous Publ NO 9504203 A; AU 9466916 A Based on WO
     9425142 A
PRAI US 1993-52395 19930423
IPCR B01D0053-46 [I,A]; B01D0053-46 [I,C];
     B01D0053-64 [I,A]; B01D0053-72 [I,A]; B01J0020-06
     [I,A]; B01J0020-06 [I,C]; C10K0001-00 [I,C]; C10K0001-20 [I,A]
AB
     WO 1994025142 A1
                        UPAB: 20050824
     The metal carbonyl content of a gas stream (I) is reduced by
     contacting the stream with lead oxide dispersed on a support.
           USE - (I) is a gas containing carbon monoxide, especially synthesis
     gas. Such a gas may form carbonyls by contact with metals, e.g. iron
     carbonyls by contact with steel processing equipment; the carbonyls
     poison catalysts in downstream conversion processes, e.g. when
     synthesis gas is used in Fischer-Tropsch, ammonia or
     methanol synthesis processes.
           ADVANTAGE - Supported lead oxide removes Fe(CO)5 rapidly from
     the gas, and has high capacity, e.g. up to 5 weight% iron on the sorbent
     trap; it is non-catalytic for synthesis and hydrogenation reactions.
ABDT W09425142
     The metal carbonyl content of a gas stream (I) is reduced by
     contacting the stream with lead oxide dispersed on a support.
     (I) is a gas containing carbon monoxide, especially synthesis gas. Such a
     gas may form carbonyls by contact with metals, e.g. iron carbonyls by
     contact with steel processing equipment; the carbonyls poison
     catalysts in downstream conversion processes, e.g. when synthesis gas
     is used in Fischer-Tropsch, ammonia or methanol synthesis
     processes.
     ADVANTAGE
     Supported lead oxide removes Fe(CO)5 rapidly from the gas, and has
     high capacity, e.g. up to 5 weight% iron on the sorbent trap; it is
     non-catalytic for synthesis and hydrogenation reactions.
     EXAMPLE
     The sorbent consisted of 21.4 weight% PbO spherical particles dispersed
     on gamma alumina of dia. 3 mm; surface area of the sorbent was 272
     m2/g, pore volume 0.42 gm3/g. Two stainless steel tubes (in parallel),
     length 2 m, dia. 25.4 mm, were filled with the sorbent, and synthesis
     gas containing 7 ppm Fe(CO)5 passed at 31.25 Nl/min., 25°C, 20
     bar, GHSV 1000, for 20 days. The amts. of iron strapped in each of 5
     zones regularly spaced along the whole length of the trap, starting at
     the inlet end, were: (1) 1.49, (2) 1.33, (3) 0.268, (4) 0.154 and (5)
     0.0042 weight%. (SJP)
     PREFERRED SORBENT
     The support is a porous refractory metal
     oxide of surface area greater than 50 m2.g, e.g. gamma alumina
     of surface area 150-300 m2/g. Lead oxide is 5-50, especially 10-30 weight% of
     the combination.
     PREFERRED PROCESS
     (I) contains at least 5 mole% carbon monoxide. It is e.g. synthesis
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USHA SHRESTHA EIC 1700 REM 4B31

gas, containing 10-90% CO, 10-90% H2 and 0-80% nitrogen. The metal carbonyl is iron, nickel or cobalt carbonyl; metal carbonyl content of

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(I) is over 5 ppm before treatment, and less than 1 ppm after
     treatment. Contact takes place at 0-100, 0-50 or 25-50°C.
FS
     CPI: E11-Q01; E11-Q02; E31-A01; E35-J; E35-U05; H04-A02; J01-E02
MC
     ANSWER 25 OF 29 WPIX COPYRIGHT 2007
L68
                                                 THE THOMSON CORP on STN
     1984-127535 [21]
                        WPIX
AN
     C1984-053889 [21]
DNC
TI
     Auto-thermal reforming by partial oxidation and steam reforming - using
     monolithic platinum-gp.-metal-containing partial oxidation catalyst
DC
     H04; H09
IN
     BUCHANAN W; FLANAGAN P; HECK R M; MCSHEA W T; YARRINGTON R M
PA
     (ENGH-C) ENGELHARD CORP; (ENGH-C) ENGELHARD MINERALS CORP
CYC
     17
     AU 8319728
                                               104[7]
PΙ
                     Α
                       19840405 (198421)* EN
     NO 8303522
                        19840424 (198423)
                     Α
     DK 8304483
                     Α
                        19840514 (198426)
                                           DA
     EP 112613
                     Α
                        19840704 (198427)
                                           EN
     JP 59097501
                     Α
                        19840605 (198428)
                                           JA
     ES 8503717
                     Α
                        19850616 (198549)
                                           ES
     CA 1210242
                     A 19860826 (198639)
     CA 1210567
                     Α
                        19860902 (198640)
     CA 1217504
                     Α
                        19870203 (198711)
                                           EN
     CA 1222631
                     A 19870609 (198727)
                                           EN
     US 4844837
                     Α
                        19890704 (198934)
                                           EN
     US 4863707
                     Α
                        19890905 (198945)
                                           EN
     US 4927857
                     Α
                        19900522 (199024)
                                           EN
     EP 112613
                     В
                        19910306 (199110)
                                           FN
     DE 3382193
                     G
                        19910411 (199116)
                                           DE
     US 5023276
                     Α
                        19910611 (199126)
                                           EN
    NO 171409
                     В
                        19921130 (199302)
                                           NO
ADT AU 8319728 A AU 1983-19728 19830929; US 4844837 A US 1982-430147
     19820930; US 4863707 A US 1982-430147 19820930; US 4927857 A US
     1982-430147 19820930; US 5023276 A US 1982-430147 19820930; US 4844837
     A US 1982-430200 19820930; US 4863707 A US 1982-430200 19820930; US
     4927857 A US 1982-430200 19820930; US 5023276 A US 1982-430200
     19820930; US 4844837 A US 1982-430320 19820930; US 4863707 A US
     1982-430320 19820930; US 4927857 A US 1982-430320 19820930; US 5023276
     A US 1982-430320 19820930; US 4844837 A US 1982-430451 19820930; US
     4927857 A US 1982-430451 19820930; US 5023276 A US 1982-430451
     19820930; US 4844837 A US 1982-430452 19820930; US 4863707 A US
     1982-430452 19820930; US 4927857 A US 1982-430452 19820930; US 5023276
     A US 1982-430452 19820930; EP 112613 A EP 1983-305887 19830929; JP
     59097501 A JP 1983-179530 19830929; NO 171409 B NO 1983-3522 19830929;
     US 4863707 A US 1989-296385 19890106; US 4927857 A US 1989-298875
     19890118; US 5023276 A US 1989-300197 19890119
    NO 171409 B Previous Publ NO 8303522 A
PRAI US 1982-430147 19820930
     US 1982-430200 19820930
    US 1982-430320 19820930
    US 1982-430451 19820930
    US 1982-430452 19820930
    US 1989-296385 19890106
    US 1989-298875 19890118
    US 1989-300197 19890119
IPCR B01J0019-24 [I,A]; B01J0019-24 [I,C]; B01J0023-00 [I,A]; B01J0023-00
     [I,C]; B01J0023-44 [I,A]; B01J0023-44 [I,C]; B01J0008-02 [I,A];
     B01J0008-02 [I,C]; C01B0003-00 [I,C]; C01B0003-00 [I,C]; C01B0003-32
     [I,A]; C01B0003-36 [I,A]; C01B0003-38 [I,A]; C01B0003-38 [I,A];
     C01B0003-40 [I,A]; C01B0003-48 [I,A]; C01C0001-00 [I,C]; C01C0001-00
```

[I,C]; C01C0001-04 [I,A]; C01C0001-04 [I,A]; C07C0001-00 [I,C]; C07C0001-04 [I,A]; C07C0027-00 [I,A]; C07C0027-00 [I,C]; C07C0027-06 [I,A]; C07C0029-00 [I,C]; C07C0029-15 [I,A]; C07C0029-151 [I,A]; C07C0031-00 [I,C]; C07C0031-04 [I,A]; C07C0067-00 [I,A]; C07C0067-00 [I,C]; C10G0035-00 [I,C]; C10G0035-02 [I,A]; C10J0003-02 [I,C]; C10J0003-16 [I,A]; C10K0003-00 [I,C]; C10K0003-02 [I,A] AU 8319728 A UPAB: 20060104

Production of synthesis gas is effected in two stages: (i) catalytic partial oxidation of a feed mixture comprising hydrocarbon feed stream, H2O and O2-containing gas; the preheated feed mixture being contacted with a monolithic (honeycomb-type) catalyst (I) comprising Pd and Pt (andopt. Rh) on a refractory metal oxide, and

(ii) catalytic steam reforming of the first-stage effluent over a Pt-Rh steam reforming catalyst.

Specifically, the feed mixture to stage (i) is controlled to give an H2O:C ratio of 0.5-5 and an O2:C ratio of 0.2-0.8, and the step is carried out at 1-142 atmospheric and at such temps. that at least part of (I) is at at least 121 deg.C above the ignition temperature of the inlet stream, providing cracking of any unoxidised 5C + hydrocarbons to light (4C or below) hydrocarbons.

The specifically claimed embodiments relate to integrated processes utilising the H2-rich product gas for the production of (A) ammonia, (B) methanol, (C) SNG and (D) liquid hydrocarbons. Very low catalytic metal loadings may be used. Operation is at relatively low H2O:C and O2:C ratios, without catalyst fouling by C deposition.

FS

AB

CPI: H04-C01; H04-C02; H04-F02C; N02-E; N02-F02

=> d 26-29 ibib abs ind

L68 ANSWER 26 OF 29 JAPIO (C) 2007 JPO on STN

ACCESSION NUMBER:

1993-168926 **JAPIO**

TITLE:

CATALYST EXCELLENT IN HEAT RESISTANCE FOR PURIFYING EXHAUST GAS OF INTERNAL COMBUSTION

ENGINE AND PRODUCTION THEREOF YAMADA SADAJI; FUNABIKI MASAKI

INVENTOR:

N E CHEMCAT CORP

PATENT ASSIGNEE(S):

PATENT INFORMATION:

PATENT	ИО	KIND	DATE	ERA	MAIN IPC
				. – – – , – – – .	
JP 0516	58926	Α	19930702	Heisei	B01J023-58

APPLICATION INFORMATION .

JP 1992-148025 STN FORMAT: 19920515 ORIGINAL: JP04148025 Heisei JP 1992-148025 19920515 -

PRIORITY APPLN. INFO.: SOURCE:

PATENT ABSTRACTS OF JAPAN (CD-ROM), Unexamined

Applications, Vol. 1993

AN 1993-168926 ·JAPIO

AB PURPOSE: To enhance capacity for keeping purifying activity by forming an active layer containing a platinum group element, activated alumina, cerium oxide, a barium compound and a zirconium compound on a support.

CONSTITUTION: Catalyst components consisting of a platinum group element, activated alumina, for example, α -alumina with a specific surface area of 10-300m<SP>2</SP>/g, cerium oxide, a barium compound such as barium hydroxide and a zirconium compound such as zirconium oxide are supported on a support having a monolithic

structure. The support is formed into a honeycomb shape from refractory metal oxide such as cordierite.

The wts. of the **platinum** group element, activated alumina, cerium oxide, the barium compound and the zirconium compound per 1L of a catalyst are respectively set to 0.02-2g, 30-200g, 10-150g, 0.1-20g (as barium oxide) and 0.1-30g.

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IC ICM B01J023-58 ICS B01D053-36

L68 ANSWER 27 OF 29 JAPIO (C) 2007 JPO on STN

ACCESSION NUMBER:

1993-115780 JAPIO

TITLE:

CATALYST FOR CLEANING EXHAUST GAS

INVENTOR:

SHIRAISHI EIICHI; BABA HIDEYUKI; TSUCHIYA KAZUO;

OHATA TOMOHISA

PATENT ASSIGNEE(S):

NIPPON SHOKUBAI CO LTD

PATENT INFORMATION:

PATENT NO KIND DATE ERA MAIN IPC

JP 05115780 A 19930514 Heisei B01J023-58

APPLICATION INFORMATION

STN FORMAT:

JP 1992-101005

19920421

ORIGINAL:

JP04101005

Heisei

PRIORITY APPLN. INFO.:

JP 1991-90624

19910422

SOURCE: PATENT

PATENT ABSTRACTS OF JAPAN (CD-ROM), Unexamined

Applications, Vol. 1993

AN 1993-115780 JAPIO

AB PURPOSE: To remove simultaneously carbon monoxide (CO), hydrocarbon (HC), and nitrogen oxides (NOx) which are harmful components contained in an exhaust gas from internal combustion engines including automobiles.

CONSTITUTION: A catalyst for cleaning exhaust gas composed of an integrated structure coated with a catalyst composition which contains (a) Pd and Rh or (b) Pd, RH, platinum as noble metals, and alkaline earth metal oxide, cerium oxide, zirconium oxide, and refractory metal oxide. A preferable

catalyst composition for one liter of the integrated structure is: 0.1-50g of alkaline earth metal oxide, 5-100g of cerium oxide, 0.1-30g of zirconium oxide. Cerium oxide and zirconium oxide are preferably in the form at least partly of complex metal oxide or solid solution. COPYRIGHT: (C)1993,JPO&Japio

IC ICM B01J023-58 ICS B01D053-36

L68 ANSWER 28 OF 29 JAPIO (C) 2007 JPO on STN

ACCESSION NUMBER:

1988-270544 JAPIO

TITLE:

INVENTOR:

PRODUCTION OF CATALYST FOR CLEANING EXHAUST GAS

FUNABIKI MASAKI; OZAKI YUKIO

PATENT ASSIGNEE(S):

NIPPON ENGERUHARUDO KK

PATENT INFORMATION:

PATENT NO KIND DATE ERA MAIN IPC

JP 63270544 A 19881108 Showa B01J023-58

APPLICATION INFORMATION

STN FORMAT:
ORIGINAL:

JP 1987-104394

19870430

JP62104394

Showa

PRIORITY APPLN. INFO.: JP 19

JP 1987-104394 19870430

SOURCE:

PATENT ABSTRACTS OF JAPAN (CD-ROM), Unexamined

Applications, Vol. 1988

AN 1988-270544 JAPIO

AB PURPOSE: To increase high temperature durability of a catalyst by preparing a slurry by adding activated alumina containing the **platinum** group element, selenium oxide and barium nitrate, etc., sticking the slurry on a carrier having an integrated structure and thereafter calcining it.

CONSTITUTION: The slurry containing activated alumina containing platinum group element, selenium oxide and barium nitrate, and/or barium formate is prepared. The catalyst for cleaning exhaust gas is produced by sticking the slurry on the carrier having the integrated structure and thereafter calcining it. Said carrier consists of refractory metal oxide or durable metal and its monolithic or three-dimensional network structure is preferable as the form of the carrier. Selenium oxide

durable metal and its monolithic or three-dimensional network structure is preferable as the form of the carrier. Selenium oxide content is preferably 10∼200g/11 catalyst obtd.

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IC ICM B01J023-58

ICS B01D053-36

L68 ANSWER 29 OF 29 JAPIO (C) 2007 JPO on STN

ACCESSION NUMBER:

1982-105240 JAPIO

TITLE:

EXHAUST GAS PURIFYING CATALYST AND PREPARATION

THEREOF

INVENTOR:

WATANABE HIROO; KAWAMATA MOTOO; YAMAKAWA KOICHI

MITSUI TOATSU CHEM INC

TOYO C C I KK

PATENT INFORMATION:

PATENT ASSIGNEE(S):

PATENT NO KIND DATE ERA MAIN IPC

JP 57105240 A 19820630 Showa B01J023-40

APPLICATION INFORMATION

STN FORMAT:

JP 1980-182020 19801224

ORIGINAL:

JP55182020 Showa

PRIORITY APPLN. INFO.: JP 1980-182020

JP 1980-182020 19801224

SOURCE:

PATENT ABSTRACTS OF JAPAN (CD-ROM), Unexamined

Applications, Vol. 1982

AN 1982-105240 JAPIO

AB PURPOSE: To provide the exhaust gas purifying catalyst which comprises materials containing each noble metal components in separate carrier layers mutually and does not generate mutual movement and mixture of each noble metal components even at high temperature and of which each catalyst components continuously develop maximum intrinsic activity respectively.

CONSTITUTION: An aqueous slurry containing one kind of a platinum group metal or a water insol. compound thereof and a refractory metal oxide (e.g.; alumina) is coated on and adhered to a carrier (e.g.; a cordierite type honeycomb carrier) and, after drying, the treated carrier is fired. Next, an aqueous slurry containing a platinum group metal or a water insol. compound thereof and a refractory metal oxide which are different from one used in the aforementioned treatment is similarly coated on and adhered to the obtained incomplete catalyst and, after drying, firing is carried out. Those treatments are repeated corresponding to a number of the platinum group metals to be desirably contained and the

objective catalyst having carrier layers containing the platinum group metals as a multilayer is completed. This catalyst perfectly purifies CO, a hydrocarbon, a combustible organic compound, NOX or the like in an exhaust gas and the exhaust gas can be made harmless.

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IC ICM B01J023-40 ICS B01J037-02

ICA B01D053-36; B01J023-46

=> d his nofile

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(FILE 'HOME' ENTERED AT 10:01:35 ON 28 AUG 2007)
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FILE 'HCAPLUS' ENTERED AT 10:01:45 ON 28 AUG 2007
L1 1 SEA ABB=ON PLU=ON US20050054524/PN
SEL RN
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FILE 'REGISTRY' ENTERED AT 10:02:00 ON 28 AUG 2007
             5 SEA ABB=ON PLU=ON (11122-73-9/BI OR 1314-62-1/BI OR
L2
               1344-28-1/BI OR 7440-06-4/BI OR 7664-41-7/BI)
             1 SEA ABB=ON PLU=ON 7664-41-7/RN
L3
             1 SEA ABB=ON PLU=ON 1344-28-1/RN
             1 SEA ABB=ON PLU=ON 1314-62-1/RN
             1 SEA ABB=ON PLU=ON 7440-06-4/RN
L6
             1 SEA ABB=ON PLU=ON 11122-73-9/RN
L7
    FILE 'HCAPLUS' ENTERED AT 10:20:08 ON 28 AUG 2007
        227949 SEA ABB=ON PLU=ON L3 OR AMMONIA
L8
         25987 SEA ABB=ON PLU=ON
                                  L5 OR VANADIA
L9
        347310 SEA ABB=ON PLU=ON L6 OR PLATINUM OR PT
L10
          3596 SEA ABB=ON PLU=ON L7 OR FECR
L11
L12
          2926 SEA ABB=ON PLU=ON
                                  L7
             1 SEA ABB=ON PLU=ON L8 AND L9 AND L10 AND L12
L13
               E REFRACTORY METAL OXIDES/CT
           325 SEA ABB=ON PLU=ON "REFRACTORY METAL OXIDES"+PFT, NT/CT
L14
            10 SEA ABB=ON PLU=ON L14 AND L8
L15
            16 SEA ABB=ON PLU=ON L8 AND REFRACTORY METAL OXIDE?
L16
L17
            16 SEA ABB=ON PLU=ON L15 OR L16
             5 SEA ABB=ON PLU=ON L17 AND L10
L18
             3 SEA ABB=ON PLU=ON L18 AND L9
L19
                                  (L17 OR L18 OR L19)
            16 SEA ABB=ON PLU=ON
L20
            16 SEA ABB=ON PLU=ON L13 OR L20
L21
             1 SEA ABB=ON PLU=ON LAYERED AMMONIA OXIDAT?
L22
          3033 SEA ABB=ON PLU=ON AMMONIA OXIDAT?
L23
             1 SEA ABB=ON PLU=ON L23 AND L14
L24
               QUE ABB=ON PLU=ON FILM? OR THINFILM? OR LAYER? OR
L25
               OVERLAY? OR OVERLAID? OR LAMIN? OR LAMEL? OR MULTILAYER?
               OR SHEET? OR LEAF? OR FOIL? OR COAT? OR VENEER? OR SHEATH?
               OR COVER?
         20003 SEA ABB=ON PLU=ON L8(L)L25
L26
             1 SEA ABB=ON PLU=ON L26 AND L14
L27
            18 SEA ABB=ON PLU=ON L26 AND L10 AND L9
L28
            15 SEA ABB=ON PLU=ON L28 AND CAT/RL
L29
               E OXIDATION CATALYSTS/CT
         91484 SEA ABB=ON PLU=ON "OXIDATION CATALYSTS"+PFT,NT/CT
L30
             5 SEA ABB=ON PLU=ON L29 AND L30
L31
            10 SEA ABB=ON PLU=ON L29 NOT L31
L32
            29 SEA ABB=ON PLU=ON L21 OR L22 OR L29 OR L31
L33
            17 SEA ABB=ON PLU=ON L33 AND AIR POLLU?/SC,SX
L34
            12 SEA ABB=ON PLU=ON L33 NOT L34
L35
             6 SEA ABB=ON PLU=ON L35 AND CAT?
L36
               QUE ABB=ON PLU=ON SUBSTRAT? OR SURFACE? OR BASE# OR
L37
               SUBSTRUCT? OR UNDERSTRUCT? OR UNDERLAY? OR FOUNDATION? OR
               PANE? OR DISK? OR DISC# OR WAFER?
           780 SEA ABB=ON PLU=ON L37 AND L12
L38
             1 SEA ABB=ON PLU=ON L38 AND L8 AND L9 AND L10
L39
             3 SEA ABB=ON PLU=ON L38 AND L8
L40
L41
             4 SEA ABB=ON PLU=ON L36 AND L37
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L42
             23 SEA ABB=ON PLU=ON L34 OR (L39 OR L40 OR L41)
     FILE 'WPIX' ENTERED AT 10:57:59 ON 28 AUG 2007
              1 SEA ABB=ON PLU=ON US20050054524/PN
30 SEA ABB=ON PLU=ON REFRACTORY METAL OXIDE?
L43
            530 SEA ABB=ON
L44
             13 SEA ABB=ON PLU=ON L44 AND AMMONIA
4 SEA ABB=ON PLU=ON L45 AND PLATINUM?
L45
            4 SEA ABB=ON
L46
L47
              1 SEA ABB=ON
                            PLU=ON L45 AND VANADIA?
L48
              4 SEA ABB=ON PLU=ON L46 OR L47
L49
             3 SEA ABB=ON PLU=ON L45 AND B01D0053?/IPC
L50
             5 SEA ABB=ON PLU=ON L48 OR L49
L51
             67 SEA ABB=ON PLU=ON L44 AND PLATINUM
            33 SEA ABB=ON PLU=ON L51 AND B01D0053?/IPC
L52
L53
             17 SEA ABB=ON PLU=ON L52 AND L37
L54
             17 SEA ABB=ON PLU=ON L53 AND CATALYST?
L55
              2 SEA ABB=ON PLU=ON L54 AND (AMMONIA OR NH3)
L56
            5 SEA ABB=ON PLU=ON L50 OR L55
     FILE 'COMPENDEX' ENTERED AT 11:05:45 ON 28 AUG 2007
L57
              O SEA ABB=ON PLU=ON L44 AND AMMONIA
     FILE 'PASCAL' ENTERED AT 11:06:40 ON 28 AUG 2007
L58
              O SEA ABB=ON PLU=ON L44 AND AMMONIA
L59
             14 SEA ABB=ON PLU=ON REFRACTORY METAL OXIDE?
L60
              O SEA ABB=ON PLU=ON L59 AND (AMMONIA OR NH3)
              O SEA ABB=ON PLU=ON L58 OR L60
L61
  FILE 'JAPIO' ENTERED AT 11:10:40 ON 28 AUG 2007
L62
             0 SEA ABB=ON PLU=ON L44 AND AMMONIA
L63
             27 SEA ABB=ON PLU=ON REFRACTORY METAL OXIDE?
L64
              O SEA ABB=ON PLU=ON L63 AND NH3
             4 SEA ABB=ON PLU=ON L63 AND PLATINUM
L66
              O SEA ABB=ON PLU=ON L63 AND VANADIA
L67
              4 SEA ABB=ON PLU=ON: L62 OR (L64 OR L65 OR L66)
     FILE 'HCAPLUS, WPIX, JAPIO' ENTERED AT 11:45:50 ON 28 AUG 2007
L68
             29 DUP REM L42 L56 L57 L61 L67 (3 DUPLICATES REMOVED)
                     ANSWERS '1-23' FROM FILE HCAPLUS
                     ANSWERS '24-25' FROM FILE WPIX
                     ANSWERS '26-29' FROM FILE JAPIO
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